



Identification of genetic loci that contribute to *Campylobacter* resistance to fowlicidin-1, a chicken host defense peptide

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Antimicrobial peptides (AMPs) are critical components of host defense limiting bacterial infections at the gastrointestinal mucosal surface. Bacterial pathogens have co-evolved with host innate immunity and developed means to counteract the effect of endogenous AMPs. However, molecular mechanisms of AMP resistance in *Campylobacter*, an important human food-borne pathogen with poultry as a major reservoir, are still largely unknown. In this study, random transposon mutagenesis and targeted site-directed mutagenesis approaches were used to identify genetic loci contributing *Campylobacter* resistance to fowlicidin-1, a chicken AMP belonging to cathelicidin family. An efficient transposon mutagenesis approach (EZ::TN™ <KAN-2> Transposome) in conjunction with a microtiter plate screening identified three mutants whose susceptibilities to fowlicidin-1 were significantly increased. Backcrossing of the transposon mutations into parent strain confirmed that the AMP-sensitive phenotype in each mutant was linked to the specific transposon insertion. Direct sequencing showed that these mutants have transposon inserted in the genes encoding two-component regulator CbrR, transporter CjaB, and putative trigger factor Tig. Genomic analysis also revealed an operon (*Cj1580c-1584c*) that is homologous to *sapABCDF*, an operon conferring resistance to AMP in other pathogens. Insertional inactivation of *Cj1583c* (*sapB*) significantly increased susceptibility of *Campylobacter* to fowlicidin-1. The *sapB* as well as *tig* and *cjaB* mutants were significantly impaired in their ability to compete with their wild-type strain 81-176 to colonize the chicken cecum. Together, this study identified four genetic loci in *Campylobacter* that will be useful for characterizing molecular basis of *Campylobacter* resistance to AMPs, a significant knowledge gap in *Campylobacter* pathogenesis.

Keywords: *Campylobacter*, fowlicidin, antimicrobial peptide, resistance

INTRODUCTION

Campylobacter species including *C. jejuni* and *C. coli* are the most common bacterial causes of human gastroenteritis in the United States and other developed countries (Allos, 2001). Human *Campylobacter* illnesses are caused primarily by *C. jejuni* (~90%) and secondarily by *C. coli* (~10%). Poultry are the major reservoir of *Campylobacter* and thus the main source for human campylobacteriosis (Friedman et al., 2000). As a commensal, *Campylobacter* could establish persistent, benign infections in chickens (up to 10¹⁰ CFU per gram of feces). To successfully colonize and persist in intestinal tracts of animals and humans, *Campylobacter* must have evolved a variety of mechanisms to counteract harsh *in vivo* conditions as well as host innate immunity. However, we know less about *Campylobacter* pathogenicity than we do about other prevalent pathogens, which impeded the development of efficient intervention strategies to prevent

and control *Campylobacter* infections in humans and animal reservoirs.

Endogenous antimicrobial peptides (AMPs) belong to the most ancient and efficient components of host defense. Defensins and cathelicidins are two major families of AMPs in humans and animals. Both defensins and cathelicidins are important defense AMPs expressed by epithelium in small intestine, consequently limiting bacterial infections at the gastrointestinal mucosal surface (Wehkamp et al., 2007). Despite the existence of a broad diversity in AMP sequences and structures, the vast majority of AMPs share a common theme in the mechanism of killing action by disruption of membrane integrity (Yeaman and Yount, 2003; Peschel and Sahl, 2006). Generally, AMPs directly interact with target cells via initial electrostatic and hydrogen bond attraction, then disrupt the structure or function of bacterial membrane by permeating lipid bilayers, forming transmembrane pore, and ultimately lead to cell death. However, transmembrane pore forming is not the only mechanism of bacterial killing by AMP. AMPs may also have intracellular targets (e.g., DNA gyrase, ribosome) and

Abbreviations: AMP, antimicrobial peptide; F1, fowlicidin-1.

AMP-mediated bacterial death may occur as a result of several independent or cooperative mechanisms of action (Yeaman and Yount, 2003). Expression of defensins and cathelicidins is either constitutive or inducible in gastrointestinal tract (Wehkamp et al., 2007). Epithelial AMPs have been observed to be induced in human intestinal epithelia upon infection by *C. jejuni* (Zilbauer et al., 2005). Recently, a full panel of chicken defensins and cathelicidins (designated as “fowlicidins”) has been identified and characterized (Lynn et al., 2004; Xiao et al., 2004, 2006; Bommineni et al., 2007; van Dijk et al., 2007). These chicken AMPs displayed potent and broad spectrum of antibacterial activity, highlighting their role in host innate defense and in the development of novel antimicrobial (peptide antibiotic) (Hancock and Sahl, 2006).

Bacterial pathogens have co-evolved with host innate defense and developed means to curtail the effect of endogenous AMPs (Yeaman and Yount, 2003; Peschel and Sahl, 2006). Different AMP resistance mechanisms have been well characterized in various enteric pathogens such as *Salmonella enterica* serovar Typhimurium and *Escherichia coli* (Yeaman and Yount, 2003; Peschel and Sahl, 2006). The best studied AMP resistance mechanisms in Gram-negative bacteria involve (1) electrostatic repulsion of AMPs by modification of cell surface, (2) proteolytic cleavage of AMPs, and (3) active extrusion of AMPs by drug efflux pumps. Other mechanisms, such as capsule production, changes in the composition of lipid fatty acids in membrane and maintenance of appropriate membrane potential, have been also implicated in AMP resistance. Covalent modification of LPS in outer membrane via two-component regulatory systems (e.g., PhoP/PhoQ), which consequently reduces the negative charge and/or membrane fluidity of LPS, has been identified as a major mechanism in these Gram-negative pathogens (Ernst et al., 2001; Yeaman and Yount, 2003; Peschel and Sahl, 2006). However, as an important strategy to evade killing by innate immunity and by potential peptide antibiotics, AMP resistance mechanisms are still largely unknown in *Campylobacter*. Recently, it has been observed that lipid A modification (Cullen and Trent, 2010; van Mourik et al., 2010) and lipooligosaccharide (LOS) production (Lin et al., 2009; Naito et al., 2010) could confer *Campylobacter* resistance to polymyxin B, an AMP produced by bacterial species *Paenibacillus polymyxa*, that has been successfully used as a model peptide to study AMP resistance in many Gram-negative pathogens. However, the polymyxin B resistance gene identified from our recent study (Lin et al., 2009) did not contribute *C. jejuni* resistance to endogenous AMPs (e.g., fowlicidins and bacteriocins), strongly suggesting that polymyxin B may not be a good surrogate for studying *Campylobacter* resistance to structurally unrelated but physiologically relevant AMPs.

In this study, we chose fowlicidin-1 (F1), a cathelicidin expressed in chicken intestine, as a model peptide to examine mechanisms of AMP resistance and identified three genetic loci (*cbrR*, *cjaB*, and *tig*) involved in F1 resistance in *C. jejuni* using an efficient *in vivo* transposon mutagenesis system. In addition, genomic analysis revealed an operon *cj1580c-1584c* that shared homology to the *sap* (sensitivity to antimicrobial peptide) operon that conferred resistance to AMP-mediated killing of different pathogens such as *Salmonella* (Parra-Lopez et al., 1993).

Site-directed mutagenesis of the *sap* operon showed that *sapB* (*cj1583c*) is involved in F1 resistance in *C. jejuni*. Chicken experiments further demonstrated that these genes play an important role in the colonization of *Campylobacter* in the intestine.

MATERIALS AND METHODS

BACTERIAL STRAINS, PLASMIDS, AND GROWTH CONDITIONS

The major bacterial strains and plasmids used in this study are listed in **Table 1**. Among the 174 *Campylobacter* isolates used for susceptibility test to chicken AMP F1, 154 isolates were *C. jejuni* and 20 isolates were *C. coli*. These *Campylobacter* strains were isolated from different hosts and geographically diverse locations described our previous study (Hoang et al., 2011a). *Campylobacter* strains were routinely grown in Mueller–Hinton (MH) broth (Difco) or MH agar at 42°C under microaerophilic conditions generated by using CampyGen Plus gas pack (Oxoid) in an enclosed jar. *E. coli* strains were grown in Luria–Bertani (LB) broth with shaking (250 rpm) or on agar at 37°C overnight. Antibiotics kanamycin (Kan) or chloramphenicol (Cm) was added in medium at a desired concentration when needed.

AMP SUSCEPTIBILITY TESTING

The susceptibilities of *C. jejuni* and *C. coli* isolates to F1 were determined by a standard microtiter broth dilution method with an inocula of 10⁶ bacterial cells/ml as described previously (Lin et al., 2002). Minimum inhibitory concentrations (MICs) were determined by the lowest concentration of specific antimicrobial showing complete inhibition of bacterial growth after 24 h of incubation at 42°C. The chicken cathelicidin F1 (>95% purity) was synthesized by Bio-Synthesis (Lewisville, TX).

In vitro SELECTION OF F1 RESISTANT *C. jejuni*

F1 was used as the selective agent to obtain spontaneous F1 resistant mutants *in vitro*. Briefly, *C. jejuni* 81–176 was grown on F1-free MH agar plates overnight. The cells were harvested and suspended in MH to final OD_{600nm} about 1.2. The cell suspensions were plated on MH agar plates containing 16, 32, or 64 µg/ml of F1. Following four days of incubation under microaerophilic conditions at 42°C, the plates were checked for the emergence of F1 resistant mutants. The F1 pre-adapted cells were also used for *in vitro* selection of F1 resistant mutants. Briefly, *C. jejuni* 81–176 F1 were grown in MH broth containing sub-lethal concentration of F1 (4 µg/ml) for five consecutive passages (2 days per passage). Following the fifth passages, cultures were centrifuged, and the pellets were suspended in MH broth to final OD_{600nm} about 1.2. The bacterial cells suspensions were also plated MH agar plates with increasing concentrations of F1 for selecting F1 resistant mutant as described above. Mutants were randomly selected, grown in MH broth, and subjected to MIC test together with the parent strain *C. jejuni* 81–176.

DNA ISOLATION AND NATURAL TRANSFORMATION

Chromosomal DNA was isolated from *Campylobacter* using the Wizard Genomic Purification Kit (Promega) according to the manufacturer’s instructions. Natural transformation (biphasic method) was performed following standard procedure (Wang and Taylor, 1990).

Table 1 | Major bacterial strains and plasmids used in this study.

| Strain or plasmid | Description | Source and reference |
|-------------------------|---|-----------------------|
| <i>C. jejuni</i> | | |
| JL241 | NCTC 11168, human isolate | Parkhill et al., 2000 |
| JL28 | 81–176, human isolate with poor colonization ability in chicken | Black et al., 1988 |
| JL242 | 81–176, human isolate with high colonization ability in chicken | Black et al., 1988 |
| S3B | <i>C. jejuni</i> strain isolated from a chicken | Hoang et al., 2011a |
| JL599 | JL28 derivative, <i>cbrR</i> ::Kan | This study |
| JL601 | JL28 derivative, <i>cjaB</i> ::Kan | This study |
| JL602 | JL28 derivative, <i>tig</i> ::Kan | This study |
| JL656 | JL242 derivative, <i>cbrR</i> ::Kan | This study |
| JL657 | JL242 derivative, <i>cjaB</i> ::Kan | This study |
| JL658 | JL242 derivative, <i>tig</i> ::Kan | This study |
| JL623 | JL241 derivative, <i>cbrR</i> ::Kan | This study |
| JL665 | JL241 derivative, <i>cjaB</i> ::Kan | This study |
| JL629 | JL241 derivative, <i>tig</i> ::Kan | This study |
| JL668 | JL656/pCbrR | This study |
| JL694 | JL657/pCjaAB | This study |
| JL695 | JL658/pTig | This study |
| JL624 | <i>C. jejuni</i> S3B derivative, <i>cbrR</i> ::Kan | This study |
| JL631 | <i>C. jejuni</i> S3B derivative, <i>tig</i> ::Kan | This study |
| JL805 | <i>C. jejuni</i> S3B derivative, <i>cjaB</i> ::Kan | This study |
| JL697 | JL242 derivative, <i>cj1583c</i> ::Cm (<i>sapB</i> ⁻) | This study |
| JL706 | JL241 derivative, <i>cj1583c</i> ::Cm (<i>sapB</i> ⁻) | This study |
| JL719 | JL242 derivative, <i>cj1584c</i> ::Cm (<i>sapA</i> ⁻) | This study |
| JL793 | JL242 derivative, <i>cj1581c</i> ::Cm (<i>saAD</i> ⁻) | This study |
| JL792 | JL242 derivative, <i>cj1580c</i> ::Cm (<i>sapF</i> ⁻) | This study |
| <i>E. coli</i> | | |
| DH5α | F ⁻ ϕ 80lacZΔM15 Δ(lacZYA-argF)U169 recA1 endA1 hsdR17 (r _k ⁻ , m _k ⁺) <i>phoA supE44 thi-1 gyrA96 relA1 λ</i> ⁻ | Invitrogen |
| JL690 | DH5α containing pCjaAB | This study |
| JL691 | DH5α containing pTig | This study |
| JL652 | DH5α containing pCbrR | This study |
| JL692 | DH5α containing pcmSapB | This study |
| JL48 | Conjugation helper strain, DH5α containing plasmid RK2013 | Lin et al., 2005 |
| PLASMIDS | | |
| pRY111 | <i>E. coli</i> - <i>C. jejuni</i> shuttle vector, Cm ^r | Yao et al., 1993 |
| pCbrR | pRY111 derivative containing 1.5 kb <i>cbrR</i> gene plus its promoter region | This study |
| pTig | pRY111 derivative containing 1.6 kb <i>tig</i> gene plus its promoter region | This study |
| pCjaAB | pRY111 derivative containing 2.7 kb <i>cjaAB</i> operon plus its promoter region | This study |
| pGEM-T Easy | PCR cloning vector, Amp ^r | Promega |
| pSapB | pGEM-T Easy containing 2.0 kb <i>Cj1583c</i> (<i>sapB</i>) gene of JL241 | This study |
| pcmSapB | pSapB with Cm resistant gene inserted in <i>Cj1583c</i> (<i>sapB</i>) gene | This study |

RANDOM TRANSPOSON MUTAGENESIS AND SCREENING OF MUTANTS WITH INCREASED SUSCEPTIBILITIES TO F1

C. jejuni 81–176 (JL28 in **Table 1**, F1 MIC = 8 μg/ml) was subjected to *in vivo* transposon mutagenesis using EZ::Tn5™ <KAN-2> Transposome (Epicentre) as detailed in our previous publication (Lin et al., 2009). Briefly, one microliter of EZ-Tn5 <KAN-2> TnP transposome complex containing 25 ng transposon was used to electroporate *C. jejuni* JL28 competent cells. The Kan^r transformants were individually picked and inoculated in 96-well microplates. Following 24 h of incubation, cultures of

mutants were replicated into microtiter plates containing 4 μg/ml of F1. Those mutants that could not grow in F1-containing media were selected from the initial plates and subjected to a second screening to confirm increased sensitivities to F1. To confirm specific genetic linkage between the transposon insertion and the increased F1 susceptibility of each mutant, backcrossing of the transposon mutations into wild-type *C. jejuni* 81–176 was performed using natural transformation. The MICs of the backcrossed mutants for F1 were determined together with parent strain. The specific transposon insertion site of each mutant was

determined by directly sequencing the genomic DNA (Lin et al., 2009). Sequence analysis was performed using DNASTar software package.

PCR AND RT-PCR

PCR was performed in a volume of 50 μ l containing 1 mM each deoxynucleoside triphosphate, 200 nM primers, 2.0 mM MgCl₂, 100 ng of *Campylobacter* genomic DNA, and 2.5 U of *Taq* DNA polymerase (Promega) or *Pfu Turbo* DNA polymerase (Stratagene). Cycling conditions varied according to the estimated annealing temperature of the primers and the expected sizes of the products (available upon request). RNA was extracted from *C. jejuni* strains using RNeasy Mini Kit (Qiagen) followed by treatment of RNA samples with RNase-free DNase (Qiagen). RT-PCR was conducted using the MasterAmp kit (Epicentre). Cycling

conditions for RT-PCR included an initial incubation at 55°C for 30 min and 94°C for 2 min followed by 40 cycles of 94°C for 15s, 52°C for 30s, and 68°C for 1 min, and a final extension at 68°C for 5 min. A RT-PCR mixture lacking the RT was included as a negative control.

COMPLEMENTATION *IN TRANS*

To complement *cjaB* mutation in isogenic CjaB mutant JL657, a 2.7-kb complete *cjaAB* operon including its 174-bp upstream and 193-bp downstream regions, was PCR amplified from NCTC 11168 using primer pairs of CjaAB-F and CjaAB-R (Table 2 and Figure 1). The PCR was performed using *pfu* DNA polymerase (Stratagene) and the blunt-ended PCR product was purified and ligated to shuttle vector pRY111 (Yao et al., 1993), which was digested with *Sma*I prior to ligation. The ligation

Table 2 | Key oligonucleotide primers used in this study.

| Primer | DNA Sequence (5'–3') | Product size (kb) ^a | Gene/Operon amplified ^a |
|--------------|---------------------------------|--------------------------------|------------------------------------|
| Tf-F | TCATGAATTCACCACTTAGCA | 1.6 | <i>Cj0193c (tig)</i> |
| Tf-R | TGCTATCATTGAAGGCAAATTTTA | | |
| CjaAB-F | TCGCCTAATGCCAAAGTTTC | 2.7 | <i>cjaAB</i> |
| CjaAB-R | TCACCATCTGCATTGCATTTA | | |
| Cj0643-F | GCAATGCGTATCAACAATCC | 1.5 | <i>cbrR</i> |
| Cj0643-R | AAAAATTTCCCTTTCTTTTGAAAAAC | | |
| Cj1583c-F | AAAAAGCCGAGGATTTGCTT | 2.0 | <i>Cj1583c (sapB)</i> |
| Cj1583c-R | CTGTGGCTATAGCATGAACGA | | |
| Cj1584c-F | CGGGTATATCTTGGCAGCAT | 1.9 | <i>Cj1584c (sapA)</i> |
| Cj1584c-R | GAAACCCCTAAGTCCCCTTTT | | |
| Cj1582c-F | CCTGTTTTGGTGCTCGTTTTA | 1.5 | <i>Cj1582c (sapC)</i> |
| Cj1582c-R | GCTCTGCATCTTGCAAACA | | |
| Cj1581c-F | CGCTTTAATTCATTGGTGTTC | 1.5 | <i>Cj1581c (sapD)</i> |
| Cj1581c-R | TTAAAATTTCCAAACCATCTTG | | |
| Cj1580c-F | TTGATCGTTTGTGGCATTCT | 1.5 | <i>Cj1580c (sapF)</i> |
| Cj1580c-R | AAAAATCAAAGCCCAAGGAAA | | |
| CmF | CGATTTAAATGCTCGGCGGTGTTCTTT | 0.8 | <i>cat</i> |
| CmR | CGATTTAAATGCGCCCTTTAGTTCCTAAAG | | |
| Sap4-F | GTG CTA AAA CGCTTA GTTTTTAGTATT | 0.6 | <i>sapB</i> |
| Sap4-R | AATCAAATGCTCTAAACGATTTAA AAA | | |
| Sap5-F | GATGCAGTG ATTAATCTTGATTT TCAGG | 0.5 | <i>sapA</i> |
| Sap5-R | TCCATTTTACAAATTTATAAGGACCTG | | |
| Cj1583c-RT-F | GGGCTTGATAAGCCTTTGCT | 0.41 | <i>Cj1583c</i> |
| Cj1583c-RT-R | AAAACGAGCACCAAACAGG | | |
| Cj1582c-RT-F | TTTTTAGCCTTGCTGCTTT | 0.40 | <i>Cj1582c</i> |
| Cj1582c-RT-R | CCACCAAGCGCCTATAAAAA | | |
| Cj1581c-RT-F | ACAAAGTGGAGTGGGCAAAA | 0.38 | <i>Cj1581c</i> |
| Cj1581c-RT-R | AATCCAAACTAGGCTCAC | | |
| Cj1580c-RT-F | GTGGTGACAGGTAAGCACA | 0.32 | <i>Cj1580c</i> |
| Cj1580c-RT-R | GGCTTCTCCTCCGCTTAACT | | |
| ClpP-RT-F | ATGATGAAGTGGCGCTTCT | 0.40 | <i>Cj0192c</i> |
| ClpP-RT-R | GCTTCTTGCTGACATGAAAA | | |
| Cj0644-RT-F | CAGGGGTGCATCCTTATGAA | 0.32 | <i>Cj0644</i> |
| Cj0644-RT-R | GCAAATGTTGCTTGCATTA | | |
| Cj0645-RT-F | ATAATTGCGAAATGGCAAG | 0.41 | <i>Cj0645</i> |
| Cj0645-RT-R | GCTTCTTGCTGACATGAAAA | | |

^aProduct sizes and amplified genes refer to those of the relevant primer pair.

mix was introduced into DH5 α by transformation. One transformant (JL690) with a plasmid bearing intact *cjaAB* operon (pCjaAB) was created. The pCjaAB from JL690 was then transferred to JL657 by tri-parental conjugation using JL48 as a helper strain (Lin et al., 2005). Similar approach was used to complement *tig* and *cbrR* mutations using primer pairs of Td-F/Td-R and Cj0643-F/Cj0643-R, respectively (Table 2 and Figure 1). The complemented strains, together with other related strains, were subjected to AMP killing assay as described below.

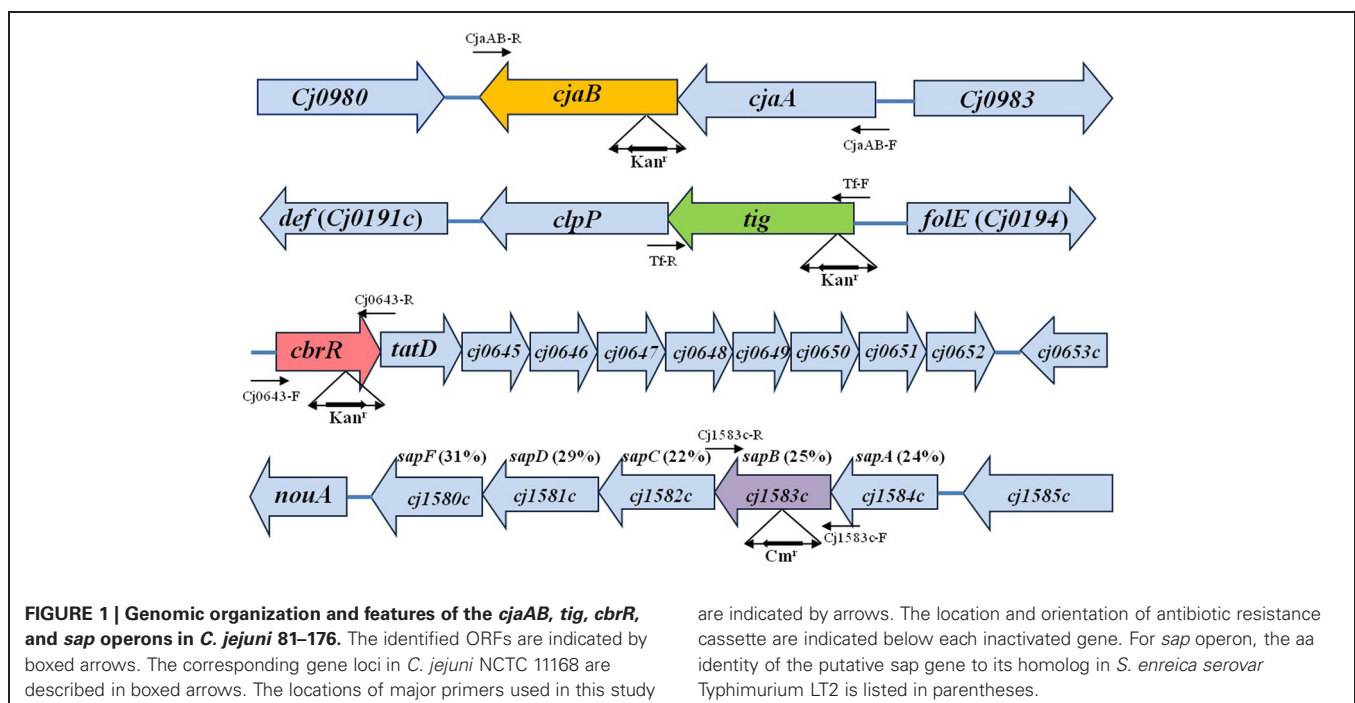
CONSTRUCTION OF ISOGENIC *sap* MUTANTS

The putative *sap* genes (*Cj1580c* to *cj1584c*, Figure 1) were inactivated by insertional mutagenesis according to our previous publication (Lin et al., 2005). Taking *sapB* (*Cj1583c*) as an example, an approximately 2 kb fragment was PCR amplified from genomic DNA of 11168 (JL241) by using primer pair Cj1583c-F and Cj1583c-R (Table 2). The PCR product was cloned into pGEM-T Easy vector (Promega) to generate pSapB (Table 1). The chloramphenicol resistance gene cassette (*cat*) was PCR amplified from plasmid pRY111 (Yao et al., 1993) by using *PfuUltra*[®] High-Fidelity DNA polymerase (Stratagene) and primers of CmF and CmR (Table 2). The resulting blunt-ended PCR product was purified and ligated into pSapB vector, which was digested with *Swa*I prior to ligation, to generate mutant construct pcm-SapB (Table 1). The construct pcmSapB, which serves as suicide vector, was then introduced into 81–176 (JL242) by natural transformation. One transformant, designated JL697, was selected on MH agar containing 5 μ g/ml of Cm. The inactivation of putative *sapB* (*Cj1583c*) in JL697 was confirmed by PCR (data not shown). A similar site-directed mutagenesis approach was used to create isogenic *Cj1584c* (*sapA*), *Cj1582c* (*sapC*), *Cj1581c* (*sapD*), and *Cj1580c* (*sapF*) mutants of 81–176 by using primers

pairs Cj1584c-F/R Cj1582c-F/R, Cj1581c-F/R, and Cj1580c-F/R, respectively (Table 2). Notably, due to the absence of a transcriptional terminator downstream of the *cat* gene and the same orientation of the *cat* gene with respect to the inactivated gene, inactivation of specific gene likely has no polar effect on the downstream genes; thus, construction of the individual isogenic *sap* mutants may help us to evaluate their relative contribution to fowlicidin resistance.

AMP KILLING ASSAY

The *C. jejuni* mutants together with their parent strains were subjected to AMP killing assay using 96-well plate as described previously (Mount et al., 2010) with minor modifications. The AMPs used in killing assay include chicken cathelicidin F1, chicken defensin AvBD9 (kindly provided by Dr. Guolong Zhang, Oklahoma State University), bacteriocin OR-7 and E-760 (Lin et al., 2009), and those purchased from Sigma (polymyxin B, colistin, cecropin A, gramicidin, and maganin). Briefly, *Campylobacter* strains were grown in MH broth to mid-log phase, and the cells were washed with MH broth and diluted to approximately 10⁶ CFU/ml in MH broth. A volume of 180 μ l of the diluted cells was mixed with 20 μ l MH broth (control) or AMP stock solution (concentration of 10-fold MIC of specific AMP) (treatment). The plates were incubated under microaerophilic conditions at 42°C for 2 h. After 2 h incubation, 20 μ l of bacterial culture were taken and serially diluted in MH broth and plated onto MH agar plates. The number of CFU was enumerated after two days of incubation under microaerophilic conditions at 42°C. Percentage survival was calculated by dividing the CFU number of bacteria incubated with AMP relative to those incubated in the presence of MH broth and then multiplied by 100. All assays were carried out in triplicate and two independent experiments were



performed. The significance of differences in susceptibility was determined using the Student's *t*-test.

CHICKEN COLONIZATION EXPERIMENT

Because *cbrR* mutant has been evaluated for its colonization ability in a previous study (Raphael et al., 2005), only *sapB* as well as *tig* and *cjaB* mutants were evaluated together with their parent strain 81–187 in this study. The chicken study was approved by the Institutional Animal Care and Use Committee at The University of Tennessee. Briefly, one-day-old broiler chickens were obtained from a commercial hatchery (Hubbard Hatchery, Pikesville, TN). The chickens were negative for *Campylobacter* as determined by culturing cloacal swabs prior to use in this study. These chickens were randomly assigned into three treatment group (10 or 11 chicks/group). At four days of age, each chicken was orally inoculated with a 1:1 mixture of wild type 81–176 (JL242) and its isogenic *cjaB* mutant (group I), *tig* mutant (group II), or *sapB* mutant (group III), with a dose of approximately 10^7 CFU of bacteria per chick. For each group, five birds were euthanized and cecal contents were collected at 3, and 10 days post-inoculation (DPI). The cecal contents from each bird were weighed and diluted in MH broth. The cecal suspensions were duplicate plated onto MH agar plates with *Campylobacter*-specific selective supplements (Oxoid, UK) for total *Campylobacter* enumeration and onto selective plates supplemented with appropriate antibiotics (30 μ g/ml of Kan or 6 μ g/ml of Cm) for the specific mutant numbers in each sample. The plating media were tested prior to use to ensure that they supported the growth of the mutant strains. Notably, before inoculation, the motility of the wild-type and its isogenic mutants were confirmed to be at a comparable level.

The number of CFU per gram of cecal contents was calculated for each chicken and was used as an indicator of the colonization level. The detection limit of the plating methods was 100 CFU/g of cecal contents. The bird from which no *Campylobacter* colonies were detected was assigned a conservative value of 99 CFU/g of cecal contents for the purpose of calculating means and for statistical analysis. Student's *t*-test was used to examine the significance of differences in *Campylobacter* colonization levels (log transformed CFU). A *P*-value of <0.01 was considered significant.

RESULTS

EMERGENCE OF ACQUIRED F1 RESISTANCE IN *Campylobacter*

Emergence of acquired AMP resistance is an important issue related to bacterial pathogenesis and development of sustainable peptide antibiotics (Yeaman and Yount, 2003; Hancock and Sahl, 2006; Peschel and Sahl, 2006). Previous studies suggested that high-level resistance to host endogenous AMP is difficult to develop (Peschel and Sahl, 2006), which is also supported by our recent *Campylobacter* work on the development of acquired resistance to polymyxin B (a widely used model peptide) and bacteriocins (the AMPs produced by bacteria) (Lin et al., 2009; Hoang et al., 2011a,b). Thus, in this study we first performed a screening to determine if high-level F1 resistance is observed in various *Campylobacter* isolates. Standard MIC test using MH broth showed that majority of *C. jejuni* and *C. coli* isolates exhibited low MIC to F1, ranging from 4 μ g/ml to 8 μ g/ml.

Among the 154 tested *C. jejuni* isolates, 58 and 85 showed MICs of 4 μ g/ml and 8 μ g/ml, respectively. Only one *C. jejuni* strain showed slightly higher MIC of 16 μ g/ml and only 6.5% of *C. jejuni* isolates (10 out of 154) showed MICs \leq 2 μ g/ml (eight isolates with MIC of 2 μ g/ml and 2 with MIC of 1 μ g/ml). *C. coli* showed similar pattern as *C. jejuni* and displayed MICs of 0.5 μ g/ml (1 isolate), 1 μ g/ml (three isolates), 2 μ g/ml (three isolates), 4 μ g/ml (eight isolates), and 8 μ g/ml (five isolates). Together, the MIC survey here showed that none of the strains displayed high-level acquired resistance to F1.

Our recent *Campylobacter* studies (Lin et al., 2009; Hoang et al., 2011a,b) and work in other bacteria have demonstrated that availability of AMP resistant mutant was very helpful for examination of acquired AMP resistance using functional genomics approaches, such as random transposon mutagenesis and microarray. Thus, in this study we also intended to obtain an *in vitro* selected F1 resistant mutant. Different *in vitro* selection methods consistently led to the emergence of F1 resistant colonies on the selective plates containing 32, or 64 μ g/ml of F1. Total 24 colonies were randomly selected for MIC test together with their parent strain 81–176. However, after being cultured in F1 free MH broth, none of these isolates displayed higher MIC than 81–176 for F1, indicating all these mutants were false-positive. Despite extensive efforts, no mutant with stable acquired F1 resistance was selected *in vitro*. In summary, these findings strongly suggest that endogenous AMPs are not favorable for the development of high-level acquired AMP resistance in *Campylobacter*.

IDENTIFICATION OF GENETIC LOCI CONTRIBUTING TO F1 RESISTANCE BY RANDOM TRANSPOSOME MUTAGENESIS

Although we failed to obtain *C. jejuni* mutants with acquired F1 resistance, examination of intrinsic AMP resistance in wild-type *C. jejuni* strains would fill a significant knowledge gap and also likely shed light on the mechanisms of acquired AMP resistance in *Campylobacter*. Thus, in this study a library containing 4800 Kan^r mutants were generated for screening the mutants with an increased susceptibility to F1. Three mutants displaying increased sensitivity to F1 were identified. Backcrossing of the transposon mutations into *C. jejuni* 81–176 further confirmed that the F1 sensitive phenotype in each mutant was linked to the gene with a specific transposon insertion. Direct sequencing of the mutant genomic DNA using transposon-specific primers showed the EZ::TN <KAN-2> transposon inserted into *Cj0981c*, *Cj0193c*, and *Cj0643* (Figure 1). All the transposon insertions occurred in the coding regions of corresponding genes. The orientations of Kan^r cassette within transposon in each mutant were the same as the corresponding disrupted gene (Figure 1). The *Cj0981c*, *Cj0193c*, and *Cj0643* encode transporter CjaB, putative trigger factor Tig, and the response regulator CbrR, respectively. F1 killing assay further confirmed that the insertional inactivation of *cbrR*, *tig*, or *cjaB* led to significantly increased susceptibility of *C. jejuni* 81–176 to F1 (Figure 2A). This phenotype was also observed in other strain background (*C. jejuni* NCTC 11168 and S3B) (Figure 2B). However, mutation of these genes did not confer significantly enhanced susceptibility of these mutants to other tested AMPs (data not shown).

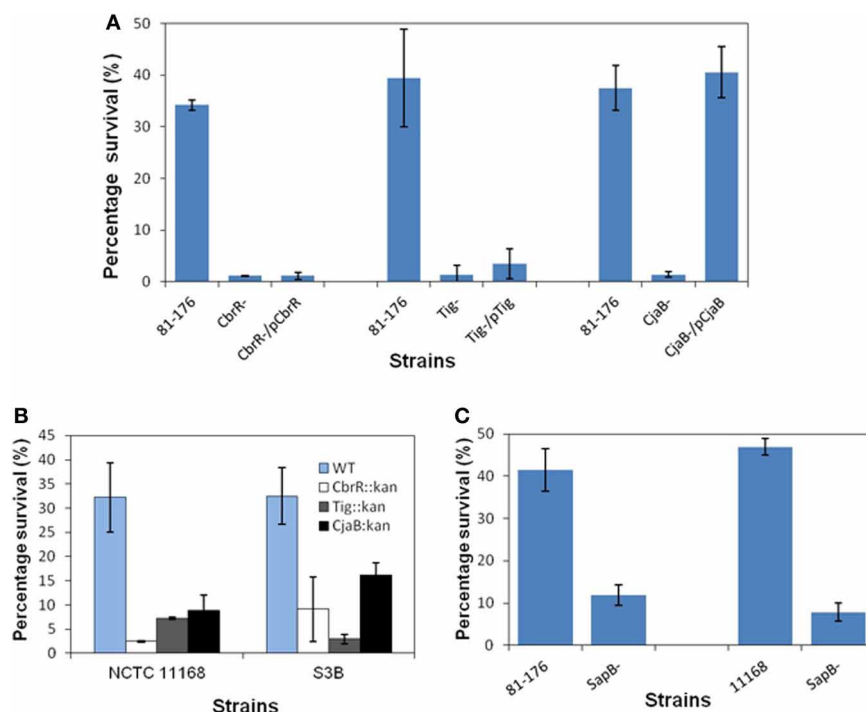


FIGURE 2 | Role of *cbrR*, *tig*, *cjaB* and *sapB* in F-1 resistance in different *C. jejuni* strains. (A) Susceptibilities of wild-type 81-176 and its *cbrR*, *tig*, and *cjaB* mutant constructs to F1. (B) Effects of mutations in *cbrR*, *tig*, and *cjaB* on the susceptibilities of *C. jejuni* NCTC 11168 and S3B to F1. (C) Inactivation of *sapB* increased susceptibilities of 81-176 and NCTC 11168 to F1. Log-phase cells in MH broth were supplemented with F1 at final concentration of 8 μ g/ml and incubated for 2 h at 42°C under

microaerophilic condition. Samples were diluted and plated on MH agar plates to determine bacterial viability. Percentage survival was calculated by dividing the CFU number of bacteria incubated with AMP relative to those incubated in the presence of MH broth and then multiplied by 100. Each data point represents the mean value obtained from two independent experiments with triplicate measurements in each independent experiment.

Complementation of *cjaB* mutation in JL657 completely restored F1 resistance level back to the level of parent strain 81-176 (Figure 2A), which is consistent with the report that the *cjaB* and its upstream gene *cjaA* formed an operon (Wyszynska et al., 2006; see Figure 1). Complementation of *CbrR* and *Tig* mutant had little effect on sensitivity of the mutants to F1 (Figure 2A), likely due to the polar effect of the transposon mutation on the downstream genes in the same operon. The *cbrR*, a gene encoding a two-component response regulator required for bile salt resistance of *Campylobacter* (Raphael et al., 2005), appears to form an operon together with its nine downstream genes (Figure 1). For example, the stop codon of *cbrR* and start codon of *tatD* overlap by 10 nucleotides (Figure 1). There is no intergenic space between the stop codon of *tig* and the start codon of downstream *clpP*, suggesting these two genes are organized into an operon. The *clpP* is separated from its downstream gene *def* by a 28 bp intergenic region which is predicted to contain promoter sequence. Thus, *tig* operon likely contains two genes (*tig* and *clpP*) (Figure 1). RT-PCR using specific primers (Table 2) showed that *clp* and *tatD*/Cj0645 were barely transcribed in *tig* and *cbrR* mutants, respectively, while all the genes were expressed normally in wild-type 81-176 (data not shown), indicating that the insertional mutation in *tig* and *cbrR* caused a polar effect on the downstream genes.

SapB CONTRIBUTES TO F1 RESISTANCE IN *C. jejuni*

The *sap* (sensitivity to antimicrobial peptide) operon confers resistance to AMP-mediated killing of different pathogens such as *Salmonella* (Parra-Lopez et al., 1993) and *Haemophilus influenza* (Mason et al., 2005) although the exact mechanisms of how Sap transporters protect the cells from AMP attack remain unclear. Analysis of NCTC 11168 genome revealed that the operon *Cj1580c-Cj1584c* shared homology to the identified *sap* operon in *S. enreica* serovar Typhimurium LT2 with identical genetic components and organization, which include (Figure 1), in which *sapA* (Cj1584c) encodes a putative periplasmic peptide binding protein and *sapB* (Cj1583c) encodes a membrane permease. Comparative genomic analysis of published *C. jejuni* genomes showed that the putative *sapABCDEF* operon is highly conserved in *C. jejuni* with nucleotide sequence identity ranging from 98 to 100%. This observation was further confirmed by our PCR survey using *sapB* specific primer pair (Sap4-F/R, Table 2) and *sapA* specific primer pair (Sap5-F/R, Table 2) for 27 diverse *Campylobacter* strains (21 *C. jejuni* strains and 6 *C. coli* isolates), in which majority of strains contain *sap* operon as reflected by the positive PCR results (23 out of 27).

Isogenic SapA, SapB, SapD, and SapF mutants of 81-176 were successfully created in *C. jejuni* 81-176 (Table 1). Mutation in SapC could not be generated after repeated attempts. RT-PCR

analysis using specific primers (Table 2) indicated that insertional inactivation using Cm resistance gene marker did not cause polar effect on the transcription of the genes (*Cj1582c*, *Cj1581c*, and *Cj1580c*) downstream of specific mutated gene (data not shown). As shown in Figure 2C, inactivation of SapB caused significantly increased sensitivity to F1 in both 81–176 and 11168 strain background. However, mutation of other individual *sap* genes including *sapA* (*Cj1584c*), *sapD* (*Cj1581c*), and *sapF* (*Cj1580c*) (Figure 1) did not alter sensitivity of the mutants to F1 (data not shown). In addition, isogenic *sapB* mutant did not display significantly increased susceptibilities to other tested AMPs with diverse sequence and structure (data not shown).

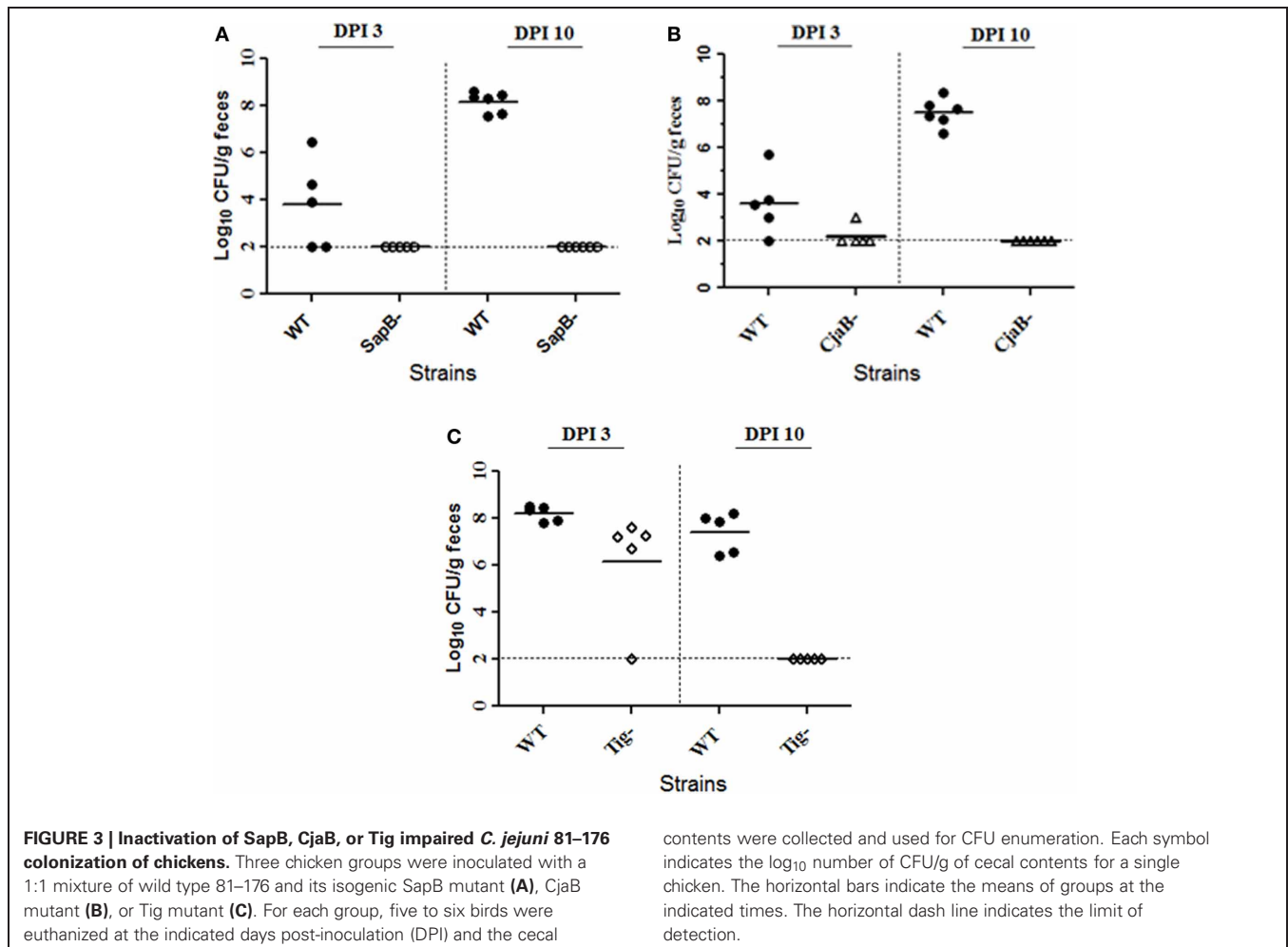
INACTIVATION OF SapB, CjaB, AND Tig REDUCED *Campylobacter* COLONIZATION IN CHICKENS

It has been demonstrated that *cbrR* is required for optimal colonization of *Campylobacter* in chickens (Raphael et al., 2005). In this study, we compared colonization ability of the isogenic *sapB*, *cjaB*, and *tig* mutants to their parent strain 81–176 using mixed infection in chicken. As shown in Figure 3, inactivation of the *sapB*, *cjaB*, or *tig* gene greatly impaired the colonization ability of *C. jejuni* 81–176 in chickens. Specifically, when 81–176 and its specific isogenic mutant were co-inoculated into a group of

chickens, the colonization level of the specific mutant was significantly lower than that of 81–176 at 3 and 10 days post-inoculation (DPI). In particular, by 10 DPI, the specific mutant of *C. jejuni* 81–176 (*tig*⁻, *cjaB*⁻, or *sapB*⁻ mutant) was no longer detected in any of the cecal samples collected from the chickens inoculated with mixture of 81–176 and its isogenic mutant. It is important to mention that both 81–176 and the isogenic mutants showed similar growth patterns in MH broth (data not shown). We also performed an identical *in vitro* competition assay in MH broth; all the mutants were not outcompeted by 81–176 and the isogenic mutant displayed an *in vitro* competitive index (ratio of the mutant to wild-type strain at early stationary phase) of about 1, indicating that growth rate of these mutants is similar to its wild-type 81–176.

DISCUSSION

Development of AMP resistance is an important strategy utilized by many bacterial pathogens including various enteric bacteria to evade host innate immunity and to colonize in various host niches successfully (Ernst et al., 2001; Yeaman and Yount, 2003; Peschel and Sahl, 2006). However, it is still largely unknown how important AMP resistance is for *Campylobacter* infection and what is the molecular basis of AMP resistance in *Campylobacter*.



To date, there are no any animal knockout experiments demonstrating the critical role of specific host AMP in defense against *Campylobacter*. However, it has been well established that AMPs (primarily defensins and cathelicidins) are a major component of host innate immunity systems against enteric pathogens. For example, mice deficient in cathelicidin or defensins production are more susceptible to intestinal infections by enteric pathogens (Wilson et al., 1999; Iimura et al., 2005). Thus, both defensins and cathelicidins are important AMPs expressed in the small intestine to protect host against pathogens (Wehkamp et al., 2007). Zilbauer et al. (2005) also observed that epithelial AMPs were induced in human intestinal epithelia upon infection by *C. jejuni*. In chickens, although different AMPs including the F1 used in this study are expressed in the intestine, the concentrations of specific AMP in any of intestinal niches are still unknown. Specific knock out model should be developed to determine if chicken deficient in F1 production is more susceptible for *Campylobacter* infection. Notably, colonization of *Campylobacter* in chicken and other hosts is determined by delicate interactions between *Campylobacter* and host in which many determinants in addition to AMP are involved. However, given the intimate interaction of *Campylobacter* with intestinal mucosa and the intracellular stage during systemic *Campylobacter* infection in humans (invasion of epithelial cells and engulfment by phagocytes), *Campylobacter* should have co-evolved with host innate defense and developed means to curtail the effect of endogenous AMPs for successful colonization as observed in other enteric pathogens (Ernst et al., 2001; Yeaman and Yount, 2003; Peschel and Sahl, 2006). However, mechanisms of AMP resistance in *Campylobacter* are still largely unknown.

We have initiated determination of AMP resistance mechanisms in *Campylobacter* using polymyxin B as a model peptide and have defined seven genetic loci (e.g., *galU* required for LOS production) contributing to *C. jejuni* resistance to polymyxin B (Lin et al., 2009). Recent work also showed that production of specific LOS (Naito et al., 2010) and *lipidA* modification (Cullen and Trent, 2010; van Mourik et al., 2010) were required for polymyxin B resistance in *C. jejuni*, likely due to their role in electrostatic shielding against polymyxin B. Polymyxin B has been successfully used as a model peptide to study AMP resistance in many Gram-negative pathogens although polymyxin B bears little structural resemblance to the host defense AMPs (Ernst et al., 2001; Yeaman and Yount, 2003; Peschel and Sahl, 2006). However, our previous study (Lin et al., 2009) strongly suggested that polymyxin B may not be a good surrogate for studying AMP resistance in *Campylobacter*. In addition, examination of *Campylobacter* resistance to bacteriocins revealed a different set of genes which are not involved in resistance to polymyxin B and other endogenous AMPs (Hoang et al., 2011a). Therefore, direct usage of natural host AMP as a model peptide may generate physiologically relevant information for *Campylobacter* pathogenesis. We chose to use F1 as a surrogate of host defense peptides because of following reasons. First, unlike defensins, the short α -helical fowlicidin-1 (F1) can be easily synthesized at reasonable cost for large scale screening. Second, cathelicidins including F1 are important AMPs expressed in intestine and displayed potent bactericidal effect (Lynn et al., 2004; Xiao et al., 2004,

2006; Peschel and Sahl, 2006; Bommineni et al., 2007). In addition, some genes (e.g., *phoPQ*) required for cathelicidin resistance could also involve resistance to structurally unrelated AMPs such as defensins (Ernst et al., 2001; Winfield and Groisman, 2004). Thus, genes contributing to F1 resistance may be important for *in vivo* colonization of *C. jejuni* in the host. Finally, F1 is produced by chicken, an animal model used in this study, which would make us well correlate the findings obtained from *in vitro* and *in vivo* studies.

In this study, inactivation of *SapB*, *CjaB*, *Tig*, and *CbrR* led to increased susceptibilities to F1 in different *C. jejuni* strain background. In addition, chicken experiment also showed impaired colonization ability of these mutants in the intestine. The *Sap* operon encode ABC-type transporter system and have been demonstrated to involve AMP resistance in various Gram-negative bacteria; mutation of the *sap* operon also resulted in impaired colonization in animal models (Parra-Lopez et al., 1993; Lopez-Solanilla et al., 1998; Chen et al., 2000; McCoy et al., 2001; Lupp et al., 2002; Mason et al., 2005, 2006; Mount et al., 2010). The exact mechanisms of how *Sap* transporters protect the cells from AMP attack remain unclear. In this study, we demonstrated the role of *sapB* in *Campylobacter* resistance to F1; however, mutation of *sapB* in *C. jejuni* did not lead to significantly increased susceptibilities to other natural AMPs. We should interpret this finding cautiously. The *C. jejuni sap* may still interact to other structurally unrelated AMPs with lower efficiency than F1; however, limited expression level of *Sap* in our *in vitro* killing assay system may diminish the susceptibility difference between *Sap* mutant and wild-type strain. Mason et al. (2003, 2005) reported that *sap* expression of *H. influenza* is greatly up-regulated *in vivo*. Therefore, the *sap* operon of *Campylobacter* is possible up-regulated during *in vivo* infection and such enhanced expression of *sap* may ultimately confer *C. jejuni* resistance to various endogenous AMPs, leading to greatly impaired intestinal colonization observed in this study (Figure 3A). This hypothesis needs to be examined in the future. In terms of *CjaAB* operon, our chicken study has clearly showed its role in *C. jejuni* colonization in the intestine, likely due to its contribution to AMP resistance. The *CjaB* was predicted as an inner membrane transporter with unknown function in *Campylobacter* (Wyszynska et al., 2006). The underlying mechanism for *CjaAB* conferring F1 resistance is not clear. It has been observed that efflux pumps are involved in AMP resistance in various bacteria (Yeaman and Yount, 2003; Peschel and Sahl, 2006). Recently, we also demonstrated that *CmeABC* efflux pump is required for *Campylobacter* resistance to bacteriocins (Hoang et al., 2011a). Therefore, it is likely that *CjaAB* function as an efflux pump to extrude F1 out of cells for resistance. However, it is equally possible that *CjaAB* may function as an influx transporter and facilitate transportation of F1 to *Campylobacter* cytoplasm where F1 is targeted for degradation.

Given the failure of complementation and lack of expression of the genes downstream of the inactivated *tig* and *cbrR*, the phenotypes of isogenic *tig* and *cbrR* mutants observed in this study are likely caused by other gene(s) in the same operon or the coordination of all genes within the operon (Figure 1). Trigger factor *Tig* was found highly conserved in eubacteria,

functioning as a chaperon to interact with newly synthesized polypeptides, assist protein folding, and plays a vital role in bacterial virulence (Rassow and Pfanner, 1996; Martinez-Hackert and Hendrickson, 2009). The gene immediately downstream of *tig* in *Campylobacter* was annotated as *clpP* that function as a protease to degrade misfolding peptides in other bacteria (Frees et al., 2007). Notably, the ClpP homolog of *Bacillus anthracis* has been demonstrated to be involved in resistance to AMPs human cathelicidin LL-3 (McGillivray et al., 2009). Therefore, since degradation of AMPs by protease is one of mechanisms used by bacteria to resist endogenous AMPs (Yeaman and Yount, 2003), the *C. jejuni* ClpP may involve proteolytic cleavage and play a role in virulence by mediating the role in AMP resistance. This hypothesis needs to be examined in the future. Recently, Cohn et al. (2007) has characterized the *C. jejuni* ClpP and observed that ClpP influenced heat tolerance of *C. jejuni*. Therefore, the impaired colonization of *tig* mutant observed in this study also could be attributed to the temperature sensitivity due to inactivation of ClpP. Similarly, *cbrR* and nine of its downstream genes appear to organize into an operon (Figure 1) and some downstream genes may be involved in AMP resistance in *C. jejuni*. For example, *Cj0649* encodes a beta-barrel LptD-like protein and the LptD in *Pseudomonas aeruginosa* appeared to act as a dominant resistance marker against cathelicidins likely due to its function in the assembly of LPS in the outer leaflet of the outer membrane (Srinivas et al., 2010). However, the two component regulators identified in this study (CbrR) and in our recent work (RacR) (Lin et al., 2009) may still involve *Campylobacter* resistance to AMPs. In response to environmental

cues via two-component regulatory systems (e.g., PhoP/PhoQ), Gram-negative bacteria can add covalent modification to LPS and consequently reduce the negative charge and/or membrane fluidity of LPS and protect themselves from attack by AMPs, which has been a major AMP resistant mechanism in enteric bacteria (Ernst et al., 2001; Yeaman and Yount, 2003; Peschel and Sahl, 2006). Mutations occurred in two-component regulators could result in sustained, acquired AMP-resistance. To hunt potential two-component regulators required for AMP resistance, we have performed sequencing of nine regulators, five histidine kinases, and six signal transduction systems in polymyxin B resistant mutant; but no sequence difference was observed in all selected regulators between wild-type 81–176 and its polymyxin B resistant derivatives (Hoang, 2010). Despite accumulating studies on the functional characterization of two-component regulators in *C. jejuni*, it is still unknown which environmental signals to which these regulators respond. We speculate that some two-component regulatory systems in *C. jejuni* can sense and integrate multiple environmental cues in the host into a coordinated cellular response and promote *Campylobacter* resistance to AMPs in different host niches. Testing this hypothesis relies on the identification of specific cues activating two-component regulatory systems, a challenge issue to elucidate physiology and pathogenesis role of two-component regulatory systems in bacterial pathogens.

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