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Comparative analysis of cytolethal distending toxin (cdt) genes, and biological activities of CDTs in *Campylobacter helveticus* and *Campylobacter upsaliensis*

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Introduction

*Campylobacter* is considered as one of the most prevalent zoonotic pathogens causing human gastroenteritis worldwide. Although the poultry and livestock animals are the main source of infection, pets are also well-recognized carriers of *Campylobacter* spp., and possess an important risk factor for human campylobacteriosis. *C. upsaliensis* and *C. helveticus* are the major campylobacters commonly found in dogs and cats, both sick and healthy animals. Considering zoonotic pathogens to humans, *C. upsaliensis* is considered to be the third predominant emerging *Campylobacter* next to *C. jejuni* and *C. coli* and campylobacteriosis in humans associated with their healthy household pets have been reported. Although information about *C. helveticus* in clinical sample is relatively rare, this species also has been detected from diarrheal patients. Thus, *C. helveticus* might be associated with human disease, which has been so far possibly underestimated. On the other hand, very little information is known about the pathogenic mechanisms and virulence factors of both *C. upsaliensis* and *C. helveticus*. Cytotoxicity, invasiveness, and persistent colonization of *C. jejuni* in vivo. Recently, *cdtB* gene has been detected from *C. helveticus* strain from healthy cat in Japan by PCR targeting a conserved region in *cdtB* genes of *C. jejuni, C. coli* and *C. fetus*. CDT-like effect was also observed in HeLa cells when cultured with the bacterial cell lysate of *C. helveticus*, and accordingly, particular strains of the bacterium might harbor entire *cdt* genes, and produce biologically active CDT. Subsequently, a *cdtB* gene-based PCR-RFLP assay was established for the detection of 7 *Campylobacter* species and this assay yielded differential RFLP patterns in *C. upsaliensis* strains isolated from healthy dogs in Japan. The objective of this study was to divulge into more details of the genetic variations among the *cdt* genes in *C. helveticus* and *C. upsaliensis*, and the biological activities of, *C. helveticus* (CheCDT) and *C. upsaliensis* CDT (CuCDT), in comparison to other campylobacters.

Chapter 1: Determination and analysis of *cdt* gene cluster in *C. helveticus* and *C. upsaliensis*

Culture followed by PCR-based screening of fecal samples from healthy cats in Japan detected the occurrence of *cdtB*-like gene in *C. helveticus* strain CAT and the gene sequence was highly homologous to the *cdtB* genes of certain *Campylobacter*. The bacterial cell lysate of the isolated *C. helveticus* strain caused cell distention in HeLa cells. To check whether this type of *C. helveticus* strain harbor the complete *cdt* genes cluster, indicating their as biologic potential, nucleotide sequences of the entire *C. helveticus cdt* (Checdt) gene cluster and its flanking region were determined and analyzed. All *C. helveticus* strains, isolated from both diarrheic and healthy cats including ATCC 51209 strain (*n*=9), were observed to harbor the complete *cdt* genes cluster with the conservation of putative amino acid residues essential for DNase I activity in the CdtB. Among genus *Campylobacter*, the deduced amino acid sequences of CheCDT were highly homologous to those of CuCDT.

With relation to the findings of a recent study showing existence of at least 3 subtypes of *Cu* *cdt* genes (RFLP patterns Cu-I, Cu-II, and Cu-III) in *C. upsaliensis* by the *cdtB* gene-based PCR-RFLP assay, it is worthy to check whether the genetic variation could be either
due to mutation in cdtB gene sequence or presence of the cdt gene variant. Therefore, in this study the entire cdt genes of 3 C. upsaliensis strains, ATCC 43954 (Cu-I), 40-1 (Cu-II) and 9-1 (Cu-III) were sequenced. Comparative genetic analysis revealed all these strains harboring cdt gene cluster with subtle differences among cdt genes of each representative strain at both nucleotide and amino acid levels. However, this of genetic variation may not be enough to consider each of the PCR-RFLP patterns representing different sub-types.

In summary, C. helveticus harbored cdt gene cluster which was found to be ubiquitously conserved among C. helveticus strains isolated from both diarrheic and healthy cats. On the other hand, C. upsaliensis analyzed in this study most likely harbored identical cdt gene variant although they showed different RFLP patterns.

Chapter 2: Evaluation of the biological activities of cdt gene-products from C. helveticus and C. upsaliensis

C. helveticus harbored the cdt gene cluster and CDT-like effect was observed in cells cultured with bacterial cell lysate of the isolate. To see if the cytotoxic effect was due to CDT, biological activities of cdt gene-products were evaluated. CDT effects including cell distention, cell cycle arrest and γH2AX were observed in cells cultured with reconstituted rCheCDT holotoxin as well as bacterial cell lysate from strain CAT. The cytotoxic effects of this strain could be neutralized by antiserum against rCheCdtA and rCheCdtC. Additionally, bacterial cell lysate of other C. helveticus strains (n=8) also caused cell distention in HeLa cells which was neutralized by the anti-rCheCdtC serum.

In case of C. upsaliensis, although Cucdt genes seem to be identical, its gene-products may possess different biological activities or may have the same. Therefore, cytotoxic effects were examined in HeLa cells with bacterial cell lysate of 3 representative strains. All strains demonstrated CDT effects which were completely neutralized by anti-rCuCdtC serum. Interestingly, CDT activity titer produced by C. upsaliensis was much higher than that of the other Campylobacters. The toxin titer among 3 representative strains is variable (1,024, 8,192 and 65,536 in Cu-II, I and III, respectively). When compared for amino acid sequences, in CdtC, arginine at position 69 was replaced with threonine in strain 99-1, showing the highest titer, when compared with the other two representative strains. Furthermore, in case of CdtA, 7 amino acid residues in CdtA of strains ATCC 43954 (Cu-I) and 99-1 (Cu-III) were conserved but differ from those in CdtA of strain 40-I (Cu-II). The difference in toxin titer could result from the difference in production level of CDT. However, the production of CdtC of strain 99-1 (Cu-III) was lowest, whereas that of ATCC 43954 was highest among these 3 strains by western blotting. It’s also possible that antiserum may have different sensitivities to CdtC from prototype strains. Western blotting with purified rCdtC protein generated from each respective strain demonstrated that the sensitivity of antiserum was identical. Interestingly, western blotting with non-denaturing protein showed that mobility of CdtC of strain 99-1, showing the highest titer, was different from that of ATCC 43954 and 40-1 even though predicted isoelectric points of holotoxin and individual toxin subunit protein did not show much difference.

In summary, C. helveticus strains produced biologically active CDT which could be the potential virulence factor of this organism, suggesting their possible emergence as important zoonotic pathogen for human. CDT activity titer produced by C. upsaliensis is
much higher than other *Campylobacters* suggesting that CuCDT might contribute more to human diseases in comparison to other CDTs. Difference in CDT activity among *C. upsaliensis* strains could be due to amino acid substitutions in CdtA and CdtC subunits as well as altered natural conformation of holotoxin, but not due to variation at the production level.

**Chapter 3: Comparative analysis of CDTs produced by pet-predominant *Campylobacters* with other CDTs in *Campylobacters***

High sequence homology was observed at amino acid level between CheCDT and CuCDT, but less to those of CjCDT and *C. hyointestinalis* CDT-II (ChCDT-II). Therefore, CheCDT and CuCDT may produce characteristics different from CjCDT and ChCDT-II. Importantly, in case of *Escherichia coli* CDT-I (EcCDT-I), transmembrane protein 181 (TMEM181) have been reported to be essential for intoxication. Therefore, the characteristics of CheCDT and CuCDT were analyzed and compared with other CDTs, in terms of cell tropism and immunogenicity. CheCDT, CuCDT, CjCDT and EcCDT-I displayed similar cell tropism. This might be due their utilization of identical toxin receptor on host cell surface. Thus, competition assay was carried out by using purified rCdtA and rCdtC, considered to be responsible for the binding of the receptors. Interestingly, rCheCdtA or rCheCdtC could competitive inhibit the activity of CheCDT and CuCDT but not CjCDT and EcCDT-I. Additionally, ChCDT-II displayed different cell tropism. On the other hand, rCuCdtC could inhibit cytotoxicity of CheCDT Neutralization assay with anti-rCheCdtA and anti-rCheCdtC sera showed inactivation of the activity of only CuCDT and vice versa. Anti-rChCdt-IIIC, anti-rCjCdtB, anti-rCfCdtB and anti-rEcCdt-IB sera evaluated in this study could not neutralize CDT activity of other CDTs except their own CDT.

In summary, CheCDT has biological characteristics related to CuCDT such as cell tropism, toxin receptor utilization and immunogenicity which are different from other CDTs in *Campylobacters*. These data indicate that CheCDT and CuCDT which are associated with pet predominant *Campylocater* have similar biological characteristics.

**Conclusions**

1. *C. helveticus* strains ubiquitously conserved *cdt* gene cluster whereas *C. upsaliensis* strains analyzed in this study harbored identical *cdt* gene variant, although exhibiting different RFLP patterns.
2. *C. helveticus* strains produced biologically active CDT which could be the potential virulence factor of this organism, suggesting their possible emergence as important zoonotic pathogen for human.
3. *C. upsaliensis* strains produced biologically active CDT and yielded much higher toxin titer than other *Campylobacters* suggesting that CuCDT might contribute more to human diseases in comparison to other CDTs. The observed difference in toxin titer among the *C. upsaliensis* strains might not be related to the production level, but due to amino acid substitutions in CdtA and CdtC, or altered natural conformation of holotoxin.
4. CheCDT and CuCDT which are produced by the *Campylobacter* species predominant among the pet animals have similar biological characteristics such as cell tropism, toxin receptor utilization and immunogenicity, which in turn were different from CDTs produced by other *Campylobacters*. 