Cholesterol Depletion Reduces Entry of Campylobacter jejuni Cytolethal Distending Toxin and Attenuates Intoxication of Host Cells

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Campylobacter jejuni is a common cause of pediatric diarrhea worldwide. Cytolethal distending toxin, produced by Campylobacter jejuni, is a putative virulence factor that induces cell cycle arrest and apoptosis in eukaryotic cells. Cellular cholesterol, a major component of lipid rafts, has a pivotal role in regulating signaling transduction and protein trafficking as well as pathogen internalization. In this study, we demonstrated that cell intoxication by Campylobacter jejuni cytolethal distending toxin is through the association of cytolethal distending toxin subunits and membrane cholesterol-rich microdomains. Cytolethal distending toxin subunits cofractionated with detergent-resistant membranes, while the distribution reduced upon the depletion of cholesterol, suggesting that cytolethal distending toxin subunits are associated with lipid rafts. The disruption of cholesterol using methyl-β-cyclodextrin not only reduced the binding activity of cytolethal distending toxin subunits on the cell membrane but also impaired their delivery and attenuated toxin-induced cell cycle arrest. Accordingly, cell intoxication by cytolethal distending toxin was restored by cholesterol replenishment. These findings suggest that membrane cholesterol plays a critical role in the Campylobacter jejuni cytolethal distending toxin-induced pathogenesis of host cells.

Campylobacter jejuni is one of the most common causative agents of food-borne infectious illnesses in humans (10, 34). Inflammatory diarrhea is commonly seen in children infected with Campylobacter species (4, 47). Infection by the pathogen in humans usually occurs through the consumption of contaminated poultry products (13). However, the virulence factors responsible for the induction of host diarrhea remain unclear.

A bacterial membrane-associated protein, cytolethal distending toxin (CDT), has been identified as one of the virulence factors required for the induction of interleukin-8 (IL-8), which is a chemokine associated with local acute inflammatory responses (20, 59). CDT is a tripartite protein toxin composed of three subunits, CdtA, CdtB, and CdtC (28), encoded by an operon comprising three subunits, CdtA, CdtB, and CdtC (28), encoded by an operon comprising cdtA, cdtB, and cdtC (46). Several bacterial species have been identified that contain CDT toxin, including Aggregatibacter actinomycetemcomitans (55), C. jejuni (22), Escherichia coli (45), Haemophilus ducreyi (12), Helicobacter hepaticus (58), and Shigella dysenteriae (41). CDT holotoxin functions as an AB2 toxin in which CdtA and CdtC form a binding (B) unit and CdtB is an active (A) unit (27). A previous study demonstrated that CdtA and CdtC can interact with the cell membrane and enable the translocation of the holotoxin across the cell membrane (38). In addition, the nuclear-translocated CdtB subunit exhibits type I DNase activity, which causes DNA damage resulting in cell cycle arrest at the G2/M phase (26).

Functional studies of CdtA and CdtC are relatively limited compared to those of CdtB. CdtA and CdtC adopt lectin-type structures that are homologous to ricin, a plant toxin (37, 38). The crystal structure of CDT from H. ducreyi revealed that it contains two important binding elements: an aromatic patch in CdtA and a deep groove at the interface of CdtA and CdtC (39). The structure-based mutagenesis study further demonstrated that mutations of the aromatic patch or groove impair toxin binding to the cell surface and reduce cell intoxication (39). The analysis of CDT from A. actinomycetemcomitans also revealed that CdtA and CdtC not only bind to the cell surface but also are associated with membrane lipid rafts (5). Lipid rafts are microdomains that contain large fractions of cholesterol, phospholipids, and glycosylphosphatidylinositol-anchored proteins (9, 21). In vitro studies showed that the structure of lipid rafts is stabilized in cold nonionic detergents such as Triton X-100 (8) but can be disrupted by the cholesterol depletion agent methyl-β-cyclodextrin (MβCD) (54). A recent study of A. actinomycetemcomitans CDT revealed that the...
CdtC subunit contains a cholesterol recognition/interaction amino acid consensus (CRAC) region, which is required for CdtC binding to cholesterol-rich microdomains (6). This finding indicates that cholesterol provides an essential ligand for Cdt binding to the cell membrane and also serves as a portal for CdtB delivery into host cells for the induction of cell intoxication.

A growing number of studies have reported that some pathogens exploit lipid rafts for toxin delivery to induce host pathogenesis (1, 5, 19, 25, 48). However, the interaction between C. jejuni CDT subunits and membrane cholesterol-rich microdomains (6). This finding indicates that cholesterol provides an essential ligand for CDT binding to the cell membrane and also serves as a portal for CdtB delivery into host cells for the induction of cell intoxication.

MATERIALS AND METHODS

Reagents and antibodies. Anti-His (His probe) and anti-proliferating cell nuclear antigen (anti-PCNA) were purchased from Santa Cruz Biotechnology (Santa Cruz, CA). Anti-caveolin-1 and anti-transferrin receptor (anti-CD71) were purchased from BD Pharmingen (San Jose, CA). Anti-actin mouse monoclonal antibodies were purchased from Upstate Biotechnology (Lake Placid, NY). Alexa Fluor 647-conjugated anti-rabbit IgG and 4',6'-diamidino-2-phenylindole (DAPI) were purchased from Molecular Probes (Invitrogen, Carlsbad, CA). ICRF-193 was purchased from Sigma-Aldrich (St. Louis, MO). M9CD, a cholesterol depletion agent which is commonly utilized to extract eukaryotic cholesterol from lipid rafts (53), was purchased from Sigma-Aldrich.

Bacterial and cell models. C. jejuni strain 7729, isolated from patients’ feces, was identified and deposited at the Chang Gung Memorial Hospital (Taoyuan, Taiwan) (57). The bacterial strain was grown on Brucella blood agar plates (Becton Dickinson, Franklin Lakes, NJ) supplemented with 10% sheep blood and 1.5% agar in a microaerophilic atmosphere at 37°C for 1 to 2 days. CHO-K1 cells (Chinese hamster ovary cells; CCL-61; American Type
Culture Collection, Manassas, VA) and AGS cells (human gastric adenocarcinoma cells; CRL-1739) were cultured in F12 medium (HyClone, Logan, UT). COLO205 cells (human colon adenocarcinoma cells; CCL-222) and Caco-2 cells (human colon adenocarcinoma cells; HTB-37) were cultured in RPMI 1640 medium (Invitrogen). All cell culture media were supplemented with 10% fetal bovine serum (FBS) (HyClone) and penicillin and streptomycin (Invitrogen).

**Construction and protein purification of CDT subunits.** Recombinant His-tagged CDT subunits were cloned by following standard protocols. DNA fragments of \( cdtA \), \( cdtB \), and \( cdtC \) were derived from the PCR amplification of \( C. jejuni \) 7729 genomic DNA. The forward and reverse oligonucleotide primers were \( cdtA-F \) (CATGCCATGGCTTGTTCTTCTAAATTTGAAAATGT) and \( cdtA-R \) (CCGCTCGAGTCGTACCTCTCCTTGGCGATATA) for the PCR amplification of the \( cdtA \) sequence, \( cdtB-F \) (CATGCCATGGCTAATTTAGAATTTTAATGTTGGC) and \( cdtB-R \) (CCGCTCGAGAAATTTTCTAAAAATTCTACTGAAAGTTGAAAGT) for the \( cdtB \) sequence, and \( cdtC-F \) (CATGCCATGGCTAATTTAGAATTTTAATGTTGGC) and \( cdtC-R \) (CCGCTCGAGAAATTTTCTAAAAATTCTACTGAAAGTTGAAAGT) for the \( cdtC \) sequence. Each \( cdt \) fragment was inserted into pET21d (Invitrogen, Carlsbad, CA) using NcoI and XhoI. Briefly, \( cdtA \) was amplified using primers \( cdtA-F \) and \( cdtA-R \) by PCR at 95°C for 10 min for 1 cycle; 35 cycles at 95°C for 1 min, 55°C for 1 min, and 72°C for 2 min; and a final extension at 72°C for 20 min. The NcoI/XhoI fragment then was ligated with pET21d to create the \( CdtA \) expression plasmid. Similar protocols were used to obtain the \( CdtB \) and \( CdtC \) expression plasmids from \( C. jejuni \) 7729 using primer pairs \( cdtB-F/ cdtB-R \) and \( cdtC-F/cdtC-R \), respectively. The PCR program used to amplify \( cdtB \) and \( cdtC \) were the same as that for \( cdtA \). The nucleotide sequence of each \( cdt \) construct was verified using the ABI Prism Dye Terminator cycle sequencing ready reaction kit (Perkin-Elmer Corp., Norwalk, CT) in an automated DNA sequencer (model 377-96; Perkin-Elmer Corp.). Sequence analysis was performed by the University of Wisconsin Genetics Computer Group (Madis...
ison, WI) package. Escherichia coli BL21(DE3) cells harboring either cdta, cdTB, or cdC expression plasmid was induced at an optical density at 600 nm (OD600) of 0.8 by 0.5 mM isopropyl β-D-thiogalactopyranoside (IPTG) at 37°C for 3 h. The expressed His-tagged CdTA, CdTB, and CdC fusion proteins were purified by metal affinity chromatography (Clontech, Palo-Alto, CA) and assessed by SDS-PAGE.

Generation of antiserum against each CDT subunit. Each purified CDT subunit (1 μg) was used to immunize a 6-week-old BALB/c mouse. All of the mice were purchased from the National Laboratory Animal Center (Taipei, Taiwan). The mice were immunized at weeks 0, 2, 4, 6, 8, 10, and 12, and the titer of the antiserum was detected at weeks 7, 9, 11, and 13. Mice were maintained in the animal center of China Medical University (Taichung, Taiwan). All procedures were performed according to the “Guide for the Care and Use of Laboratory Animals” (National Research Council) and were approved by the animal experiment committee of China Medical University (Taichung, Taiwan). The titers of antibodies against the CDT subunits in the serum were determined by enzyme-linked immunosorbent assay (ELISA). Ninety-six-well plates were coated with 500 ng of purified recombinant CDT subunits and blocked with 2% bovine serum albumin (BSA) in TBS (0.1 M Tris-HCl, pH 7.5, 0.03 M NaCl). Serial dilutions of the antiserum (1:1,000, 1:2,000, 1:4,000, 1:8,000, and 1:16,000) in TBS-Tween 20 were added to each well. Bound antibody was detected by horseradish peroxidase (HRP)-conjugated secondary antibodies (Invitrogen) and quantified by measuring the optical density at 450 nm after development with the TMB substrate buffer system (Kirkegaard & Perry Laboratories, Gaithersburg, MD).

SDS-PAGE and Western blot analysis. CDT holotoxin-treated cells were washed three times with phosphate-buffered saline (PBS) and then boiled in SDS-PAGE sample buffer for 5 min. The samples then were resolved by 12% SDS-PAGE and transferred onto polyvinylidene difluoride membranes (Millipore, Billerica, MA). The membranes were incubated with antiserum against each CDT subunit and then incubated with HRP-conjugated secondary antibodies (Invitrogen). The proteins of interest were visualized using the ECL Western blotting detection reagents (GE Healthcare, Piscataway, NJ) and detected using X-ray film (Kodak, Rochester, NY).

Cytoloidal distending phenotype. CHO-K1 cells were cultured at 37°C for 20 h in six-well plates containing F12 medium supplemented with 10% FBS. After one wash with PBS, cells were exposed to an individual recombinant CDT subunit (200 nM) or CDT holotoxin (200 nM each subunit) for 48 h. The CDT-treated cells were observed using an inverted optical microscope (Carl Zeiss, Göttingen, Germany).

Flow cytometry analysis of cell cycle. CHO-K1 cells treated with CDT holotoxin were analyzed by flow cytometry. Cells were pretreated with MβCD (Sigma-Aldrich) for 1 h, washed, and exposed to CDT holotoxin or an individual CDT subunit for an additional 48 h. Cells were harvested and fixed with ice-cold 70% ethanol for 1 h and stained with 20 μg/ml propidium iodide (Sigma-Aldrich) containing 1 mg/ml RNase (Sigma-Aldrich) for 1 h. The stained cells were analyzed with an FACSCalibur flow cytometer (Becton Dickinson, San Jose, CA). The data were collected using 20,000 cells from each sample and analyzed using Cell Quest software WinMDI (Verity Software House, Topsham, ME). All experiments were performed in triplicate from at least three independent experiments; all data are representations of typical experimental outcomes.

Detection of cellular cholesterol and cell viability assay. To measure the cholesterol levels in total cell lysates or detergent-resistant membrane (DRM), CHO-K1 cells were treated with various concentrations of MβCD. After incubation at 37°C for the indicated periods, the treated cells were washed three times with PBS and then disrupted by ultrasonication (three 10-s bursts at room temperature). The cholesterol content then was measured using an Amplex Red cholesterol assay kit (Molecular Probes, Eugene, OR). The percentage of remaining cholesterol after pretreatment with MβCD was determined as (fluorescence of treated cells obtained from a standard curve/total fluorescence of untreated cells) × 100 as previously described (25). To test the influence of MβCD on cell viability, cells were incubated with various concentrations of MβCD at 37°C for 1 h. After that, cells were washed three times with PBS and then incubated with fresh medium containing 10 μM lovastatin (Sigma-Aldrich) to inhibit cellular cholesterol biosynthesis. After incubation for a further 24 h, the viability of the cells was determined by using a trypan blue exclusion assay. In brief, equal volumes of 0.2% trypan blue (Sigma-Aldrich) and cell suspension were mixed. A volume of 10 μl of the mixture was placed on the hemocytometer for counting trypan blue-stained cells. A total of 300 cells in randomly selected fields were counted by a light microscope. The percentages of live and dead cells were calculated as cell viability (percent) = (live cell count/total cell count) × 100. The analysis was examined in three independent studies, each conducted in duplicate.

FIG. 4. Cholesterol depletion in CHO-K1 cells by treatment with MβCD. (A) CHO-K1 cells were treated with 10 mM MβCD at 37°C and incubated for the indicated times. The cells were harvested and subjected to cold detergent extraction using 1% Triton X-100, followed by centrifugation to isolate the DRM fraction. The prepared total cell lysates and DRM fraction then were analyzed for cholesterol concentration as described in Materials and Methods. (B) CHO-K1 cells were treated with various concentrations of MβCD (0, 2.5, 5, 10, and 20 mM) for 1 h. Whole-cell lysates and the DRM fraction then were prepared for cholesterol level analysis. (C) Cell viability was barely influenced after treatment with 0 to 20 mM MβCD, as determined by the trypan blue exclusion assay. The data represent the means and standard deviations from three independent experiments. An asterisk indicates P < 0.05 compared to results for each untreated control group, as determined by Student’s t test. DRM, detergent-resistant membrane; MβCD, methyl-β-cyclodextrin.

Immunofluorescence labeling of CDT-treated cells. To visualize the localization of CDT in cells, CHO-K1 cells (0.5 × 10⁴) were seeded on coverslips in six-well plates and incubated for 20 h. Cells were cultured with an individual CDT subunit (200 nM) or CDT holotoxin (200 nM each subunit) at 11°C for 1 h to maintain the fluidity of cell membrane and to prevent the internalization of cells (24). The cultured cells then were washed three times with PBS and fixed with 3.7% paraformaldehyde (Sigma-Aldrich) for 1 h. The cells were permeabilized with 0.1% Triton X-100 for 30 min and stained with anti-caveolin-1 antibody (BD Pharmingen), followed by being stained with Alexa Fluor 647-conju-
gated anti-rabbit IgG (Molecular Probes). To label the individual CDT subunit, samples were incubated for 30 min with anti-CdtA, anti-CdtB, or anti-CdtC antiserum followed by fluorescein isothiocyanate (FITC)-conjugated AffiniPure goat anti-mouse IgG (Jackson ImmunoResearch, West Grove, PA). Samples were analyzed under a confocal laser-scanning microscope (Zeiss LSM 510; Carl Zeiss, Göttingen, Germany) with a 100× objective (oil immersion; aperture, 1.3).

The distribution of fluorescence intensity for each CDT subunit and caveolin-1 was analyzed by ZEN software (Carl Zeiss) and schemed as line intensity histograms.

Isolation and analysis of DRM fraction. To isolate detergent-soluble and detergent-resistant fractions, CDT-treated cells were lysed with ice-cold TNE buffer (25 mM Tris-HCl, pH 7.5, 150 mM NaCl, 5 mM EDTA) containing 1% Triton X-100 and incubated on ice for 30 min. Cell lysates were centrifuged at 18,000 × g at 4°C for 30 min to separate the detergent-soluble and detergent-resistant fractions as described previously (53). Each fraction then was analyzed by Western blotting.

Isolation of nuclear fractions. To study the localization of CdtB in the nucleus of target cells, CHO-K1 cells were incubated in the presence or absence of CdtB.
absence of 10 mM MβCD at 37°C. After 1 h, cells were exposed to CDT holotoxin at 37°C for the indicated periods. The nuclear proteins then were prepared using a nuclear extraction kit (Pierce, Rockford, IL). All protein concentrations were determined by colorimetric assay using the Bio-Rad assay kit (Bio-Rad, Hercules, CA). The isolated proteins (20 μg) from the nuclear fractions then were subjected to Western blotting for further analysis of CdtB localization.

Statistical analysis. The Student’s t test was used to calculate the statistical significance of experimental results between two groups. P < 0.05 was considered statistically significant.

Nucleotide sequence accession numbers. The nucleotide sequences of cdta and cdtc have been submitted to GenBank under accession numbers JF520784 and JF682840, respectively.

RESULTS

Expression and functional analysis of recombinant C. jejuni CDT subunits. We first investigated the activity of recombinant C. jejuni CDT using Chinese hamster ovary (CHO-K1) cells. Each C. jejuni CDT subunit was cloned and expressed with a His tag in E. coli BL21(DE3) cells. Recombinant CDT subunits then were purified and analyzed by SDS-PAGE (Fig. 1A). The purified recombinant CDT subunits were readily detected by Western blot analysis using a monoclonal anti-His antibody (Fig. 1B). Western blotting was carried out to determine whether polyclonal antibodies generated against each subunit could recognize the individual CDT subunits when they were assembled and associated with cells. As shown in Fig. 1C, the individual recombinant CDT proteins were recognized by the respective polyclonal CDT antiserum (anti-CdtA, anti-CdtB, or anti-CdtC). Thus, the polyclonal antisera were further applied to investigate the association of the CDT subunits with cell membrane lipid rafts. To characterize the biological function of C. jejuni CDT holotoxin, we examined its ability to induce cell distention in CHO-K1 cells. Our results revealed that no individual recombinant CDT protein had any effect on the cell cycle or the morphology of CHO-K1 cells after culture for 48 h (Fig. 1D). However, upon the exposure of the cells to CDT holotoxin (200 nM each subunit) for 48 h, cell cycle analysis showed G2/M arrest, and light microscopy indicated cell distention (Fig. 2).

To test whether CDT has the ability to intoxicate other cell types, we employed three different cell lines (AGS, CHO-K1, and COLO205 cells) to determine the intoxication activity of CDT. Each cell line was treated with various concentrations of CDT, and the cell cycle distribution was analyzed by flow cytometry. As shown in Fig. 3A, CDT-induced cell cycle arrest at the G2/M phase in these three cell lines occurred in a dose-dependent manner. The saturation dose of CDT in AGS was ~200 nM. In contrast, the saturation doses of CDT in CHO-K1 and COLO205 cell lines were even higher, sometimes higher than 500 nM. It appeared that AGS cells were more sensitive to CDT than the other two cell lines. We then determined the binding activity of each CDT subunit to CHO-K1 cells by flow cytometry. CdtA bound to CHO-K1 cells at a saturation level of ~10 nM (Fig. 3B). The binding activity of CdtC was higher than that of CdtA at a saturation concentration of ~200 nM; however, CdtB showed no detectable binding activity to CHO-K1 cells. These results indicated that CdtA and CdtC have binding activity to CHO-K1 cells while CdtB did not.

Cholesterol is required for the association of C. jejuni CDT with the cell membrane. To determine whether cholesterol is important for the association of the C. jejuni CDT subunits with the membrane, we evaluated the ability of MβCD to deplete cholesterol from cells and membrane rafts (also called the detergent-resistant membrane [DRM]). As shown in Fig. 4A, the cholesterol concentration in total cell lysates and the DRM was decreased as early as 10 min after the treatment of CHO-K1 cells with 10 mM MβCD. Furthermore, the cholesterol levels of total cell lysates and the DRM were reduced in a dose-dependent manner by MβCD treatment for 1 h. The result showed that more than 60 and 90% of the cellular and DRM cholesterol, respectively, was extracted when cells were treated with 10 mM MβCD (Fig. 4B). We then examined the cytotoxic effect of various concentrations of MβCD on CHO-K1 cells; the cells remained essentially viable even when they were treated with 20 mM MβCD for 1 h followed by incubation with lovastatin for a further 24 h (Fig. 4C).

We then assessed the association of the CDT subunits with membrane rafts. CHO-K1 cells first were incubated with 200 nM the individual CDT subunits for 2 h at 4°C. Subsequently, the cells were analyzed by flow cytometry for the presence of CDT proteins on the cell membrane. As shown in Fig. 5 (upper), CdtA and CdtC were associated with the cell membrane. The mean channel fluorescence levels (MCF) for anti-CdtA and anti-CdtC antibodies were 42.4 and 130.0, respectively; however, the MCF for anti-CdtB antibody was only 11.0. When CHO-K1 cells were pretreated with 10 mM MβCD for 1 h, the MCF were reduced for both anti-CdtA (26.4) and anti-CdtC (32.1) antibodies (Fig. 5, lower); however, the MCF of the anti-CdtB (10.2) antibody remained similar to the level of cells not treated with MβCD. We then examined whether cholesterol depletion could affect the binding of CDT holotoxin to
cells. Noticeably, compared to the level for the control CHO-K1 cells (Fig. 6, upper), the pretreatment of CHO-K1 cells with M9252CD reduced CDT holotoxin binding (Fig. 6, lower). Taken together, these results indicate that both CdtA and CdtC are the key subunits binding to DRM, and the activity of CdtB in holotoxin depends on the binding activity of CdtA and CdtC.

We next analyzed the detergent solubility of membranes from CDT-treated cells to determine whether recombinant C. jejuni CDT holotoxin could associate with lipid rafts. CHO-K1 cells were exposed to CDT holotoxin (200 nM each subunit) for 2 h at 37°C. The cells then were collected and treated with ice-cold 1% Triton X-100 for 30 min, followed by gentle centrifugation to separate the DRM and soluble fractions. Western blotting showed that a raft-associated protein, caveolin-1, was enriched in the DRM fraction (Fig. 7). In contrast, a non-raft-associated protein, CD71 (also known as the transferrin receptor), was distributed mainly in the soluble fraction. The localization of CdtA and CdtC were enriched in the DRM fraction of CHO-K1 cells but not in the soluble fraction (Fig. 7). When cells were exposed to CDT holotoxin, the majority of CdtB was associated with DRM fraction, but a small portion of this protein was found in the soluble fraction. However, when cells were incubated with CdtB alone, CdtB was not detected in the

FIG. 8. Association of CdtA and CdtC with membrane rafts. CHO-K1 cells were exposed to medium alone or 200 nM each recombinant C. jejuni CDT subunit at 11°C for 1 h. The cells were washed and treated with control preimmune serum or each anti-CDT antiserum and then probed with FITC-conjugated anti-mouse IgG (green). The cells were co-stained with anti-caveolin-1 and Alexa Fluor 647-conjugated anti-rabbit IgG to visualize the raft microdomains (red) and analyzed by confocal microscopy. The colocalization of each CDT subunit with lipid raft domains appears as yellow in the overlay. Scale bars, 10 μm. The distribution of fluorescence intensity for each CDT subunit and Cav-1 signals across the blue lines were calculated and are presented as line intensity histograms in the right panels. Cav-1, caveolin-1.
DRM or soluble fractions (data not shown). These results were consistent with the data presented in Fig. 5 and 6, suggesting that the delivery of \textit{C. jejuni} CdtB requires the association of both CdtA and CdtC subunits with the membrane rafts in target cells.

**Delivery of CdtB into cells requires the association of CdtA and CdtC with raft microdomains.** We next used confocal microscopy to visualize whether the distribution of \textit{C. jejuni} CDT subunits is raft associated. CHO-K1 cells were incubated with 200 nM of the individual CDT subunits at 11°C for 1 h to maintain cell membrane fluidity and prevent internalization. The cells then were stained with preimmune serum and antisera to CdtA, CdtB, or CdtC. The cells were exposed to an anti-caveolin-1 antibody to identify the membrane raft microdomains. As expected, there was no signal for CDT subunits in untreated CHO-K1 cells, whereas caveolin-1 staining was observed around the plasma membrane (Fig. 8, first row). When cells were treated with each of the CDT subunits at 11°C for 1 h, CdtA (green in Fig. 8, second row) and CdtC (green in Fig. 8, fourth row) colocalized with caveolin-1 (red) on the plasma membrane. In contrast, no CdtB fluorescence could be detected on the plasma membrane (Fig. 8, third row). We then examined CDT holotoxin with confocal microscopy. Imaging analyses showed that CdtA (Fig. 9A) and CdtC (Fig. 9C) colocalized with caveolin-1 (red). In parallel, CdtB was associated with membrane rafts (Fig. 9B). The analysis of the distribution of fluorescence showed that all three CDT subunits colocalized with the membrane raft marker caveolin-1 (Fig. 9D, E, and F). These data were consistent with our findings for CDT binding activity using flow cytometry (Fig. 5 and 6), indicating that CdtA and CdtC not only associate with the membrane but also colocalize with the cholesterol-rich microdomains.

To further test whether CdtB transport to the host cells is dependent on the association of CdtA and CdtC with membrane lipid rafts, cells were exposed to CDT holotoxin (200 nM each subunit) at 37°C for 1 to 6 h. As expected, we first observed the cytoplasmic distribution of CdtB (green) at 2 h and then a clear nuclear distribution in cells without MβCD (Fig. 10A, upper). However, the amount of CdtB-associated fluorescence detected at the nucleus was visibly reduced when the cells were pretreated with 10 mM MβCD (Fig. 10A, lower). We then examined whether the nuclear localization of CdtB was dependent on the presence of cholesterol. The cells were pretreated with 10 mM MβCD for 1 h and then exposed to CDT holotoxin for 1 to 6 h at 37°C. As shown in Fig. 10B and C, the nuclear localization of CdtB gradually increased with incubation time in control cells, but its nuclear localization was almost completely blocked in cells with MβCD treatment. Taken together, these observations support the notion that the binding of \textit{C. jejuni} CdtA and CdtC to lipid rafts is important for the delivery of CdtB to target cells.

**Cholesterol depletion prevents CDT-induced cell cycle arrest.** To determine whether \textit{C. jejuni} CDT-induced cell cycle arrest depended on membrane rafts, we investigated whether the integrity of cholesterol-rich microdomains is essential for CDT holotoxin-induced cell cycle arrest. Only 17% of CHO-K1 control cells were in G2/M phase, reflecting normal

![Fig. 9. Localization of CdtB in membrane rafts through the association of CdtA and CdtB with rafts. CHO-K1 cells were exposed to 200 nM CDT holotoxin for 1 h at 11°C. The cells were washed and stained with each anti-CDT antiserum and then probed with FITC-conjugated anti-mouse IgG (green). The cells were costained with anti-caveolin-1 and Alexa Fluor 647-conjugated anti-rabbit IgG to visualize the raft microdomains (red). The stained cells were then analyzed using a confocal microscope. Scale bars, 10 μm. The distribution of fluorescence intensity for individual CDT subunits and Cave-1 signals across the blue lines was calculated and presented as line intensity histograms in the lower panels. Cave-1, caveolin-1.](image-url)
cell cycle distribution (Fig. 11A). Cells incubated with 5 or 10 mM MβCD at 37°C for 1 h did not alter the cell cycle distribution, similarly to the control cells (Fig. 11B and C). In the presence of 2 μg/ml ICRF-193, a DNA topoisomerase II inhibitor (3), 60% of cells were accumulated in G2/M (Fig. 11D), which was used as a positive control for a typical cell cycle arrest.

After pretreating CHO-K1 cells with 0, 5, or 10 mM MβCD at 37°C for 1 h, the cells were incubated with CDT holotoxin for 48 h after the removal of MβCD, and the number of cells arrested in G2/M clearly decreased in a dose-dependent manner (Fig. 11E to G). Apparently, cholesterol depletion by MβCD, which disrupts the integrity of rafts, also diminishes the activity of CDT, leading to the reduction of G2/M arrest. Upon the replenishment of cholesterol, the inhibitory effect of MβCD on CDT-induced G2/M arrest was reversed (Fig. 11H and I). Taken together, these results indicate that the presence of sufficient cholesterol in membrane raft microdomains is required for the activity of C. jejuni CDT.

We further analyzed whether the depletion of cholesterol affects CDT-induced cell cycle arrest at the G2/M phase in other cell types. Three intestinal-derived cell lines (AGS, COLO205, and Caco-2 cells) were employed in this study. Cells were untreated or were pretreated with MβCD for 1 h and then exposed to CDT holotoxin (200 nM) for 48 h. When cells were incubated with CDT holotoxin, levels of cell cycle arrest at G2/M were significantly higher in CHO-K1, AGS, COLO205, and Caco-2 cells (Fig. 12A). The number of cells arrested in G2/M was significantly decreased in CHO-K1, AGS, COLO205, and Caco-2 cells upon the pretreatment of cells with MβCD. We further assessed whether the depletion of membrane cholesterol affects the association of the CDT subunits with cell surface. Cells were incubated with 200 nM each CDT subunit and analyzed by flow cytometry for the binding of CDT proteins on the cell membrane. As shown in Fig. 12B, CdtA and CdtC were associated with the cell membrane in all tested cell lines, but CdtB was not. After pretreating cells with MβCD, the binding activity of CdtA and CdtC was reduced significantly in all tested cells. These results indicated that cholesterol is important for the association and intoxication of C. jejuni CDT in host cells. Apparently, this...
effect is observed not only in CHO-K1 cells but also in other cells.

**DISCUSSION**

This study provides evidence that the interaction of *C. jejuni* CdtA and CdtC with cholesterol-rich membrane microdomains is essential for the delivery of CdtB to target cells. The association of *C. jejuni* CDT subunits with membrane lipid rafts was investigated using flow cytometry and biochemical analyses. Our data indicated that both the CdtA and CdtC subunits, but not the CdtB subunit, are capable of binding to the cell membrane, and this binding activity was reduced after cholesterol depletion by MβCD. The CdtB subunit alone neither bound to the cell surface nor associated with lipid rafts; in contrast, CdtB was identified in the DRM fraction from cells incubated with CDT holotoxin. These findings are consistent with the crystal structure of CDT from *H. ducreyi* (38, 39). CdtA and CdtC, which have high homology to ricin proteins, have been shown to have prominent molecular surfaces, an aromatic cluster, and a deep groove that contribute to their membrane interactions (38). Another report indicated that the three CDT subunits from *A. actinomycetemcomitans* form a functional toxin unit on the cell surface which requires complex formation between the CdtA and CdtC subunits (51). These results indicate that the three subunits need to be assembled prior to holotoxin binding to the cell membrane. This is consistent with our observation that, in the presence of holotoxin, *C. jejuni* CdtB was detected predominantly in the DRM fraction of target cells, and this interaction likely is mediated through the association of CdtA and CdtB with lipid rafts.

In this study, we first employed CHO-K1 to study the mechanism of action of CDT holotoxin. Many studies also used this model for the analysis of CDT functions in *Campylobacter* spp. (2, 22, 36, 40), *A. actinomycetemcomitans* (11, 31, 32), *Escherichia coli* (7, 43), and *Shigella dysenteriae* (41) for decades. Thus, it appears that CHO-K1 is a good model for delineating mechanisms of CDT. Nevertheless, we further employed three additional intestinal-derived cell lines (AGS, COLO205, and Caco-2 cells) to determine whether cholesterol plays a crucial role in CDT binding and its activity. Our results conclude that the depletion of cholesterol affects CDT function not only in CHO-K1 cells but also in other cell lines (Fig. 12).

The CRAC region contains the conserved motif L/V(X)1-5Y(X)1-5R/K, which is present in proteins that associate with cholesterol (29). A recent report showed that the CdtC subunit of *A. actinomycetemcomitans* contains a CRAC region, which may contribute to the interaction between CdtC and cholesterol (6). Our data and another study (5) indicated that CdtA and CdtC were localized mainly in the cholesterol-rich microlayer.
We then analyzed the conserved region within the C. jejuni CdtC subunit (44), i.e., the amino acid sequence that represents a CRAC-like region (77LPFGYVQFTNPK88) (Fig. 13). C. jejuni CdtA also possesses a conserved CRAC-like motif (17LYACSSK23). Thus, these observations indicate that C. jejuni CdtA and CdtC contain hypothetical CRAC regions that contribute to their cholesterol-binding activity. However, we did not demonstrate a direct interaction between CdtA/C and cholesterol. Moreover, not all proteins that bind cholesterol have CRAC domains. A recent report indicated that only two amino acids are responsible for the recognition of cholesterol by cytolysin (16). Certainly, further investigation is needed to determine whether the CRAC-like motifs are the most important regions for the interaction of C. jejuni CdtA and CdtC with cholesterol.

The importance of cellular cholesterol for another CDT from A. actinomycetemcomitans has been well documented by Boesze-Battaglia and colleagues (5). These authors used either confocal microscopy or flow cytometry analysis. Similarly, we also found that CdtA/C of C. jejuni could bind to the cell membrane, particularly in the raft microdomains, and this effect was responsible for its toxicity activity. However, we notice some differences of CDT between A. actinomycetemcomitans and C. jejuni from this study. For example, CDT from A. actinomycetemcomitans has been demonstrated to interact with the glycosphingolipids GM1, GM2, and GM3 (35). CHO-K1 cells lack GM2 synthase, which is an upstream enzyme required for GM1 synthesis (49), suggesting that the binding of CDT from C. jejuni to the receptors in the cholesterol-enriched microdomains is different from that of A. actinomycetemcomitans. Noticeably, in this study we employed cell models resembling the natural host for C. jejuni. Therefore, we believe that the outcome of this study reflects the physiological relevance of this toxin.

Several studies have reported that lipid rafts might serve as an entry site for pathogens. For instance, in Shigella flexneri, the bacterial invasin IpaB interacts with raft-associated CD44 within specialized membrane microdomains (24). Type 1 fimbriated E. coli also was found to be associated with caveolae and lipid raft components (14). Bacteria hijack lipid rafts to mediate the infectious process; similarly, lipid rafts are required for the translocation of cytotoxin-associated gene A proteins.
(CagA) as well as for the delivery of vacuolating cytotoxin (VacA) into host cells following H. pylori infection (17, 23, 25, 50). These results suggest that lipid rafts are not only a dynamic structure on the cell membrane but also provide a bacterial entry site for toxin delivery into target cells. Our study suggests that the association of CdtA and CdtC with membrane rafts mediates the action of the toxin more efficiently. This idea is supported by the finding that the A. actinomycetemcomitans CdtB subunit exhibits phosphatidylinositol-3,4,5-triphosphate (PIP3) phosphatase activity similar to that of phosphatase and tensin homolog deleted on chromosome 10 (PTEN) (30, 52). A previous study found that PTEN normally has a generalized cytosolic and membrane distribution but is recruited into membrane rafts when cells are treated with ceramide (18). We found that both CdtA and CdtC interact with lipid rafts (predominantly localized in cholesterol-rich microdomains) to enhance the association of CdtB with the cell membrane and its subsequent delivery into the cells. This association may be important for the toxin to hijack lipid rafts for the regulation of PIP3 signaling and thereby increase the efficiency of cell intoxication.

In a recent report, Eshraghi et al. presented a comprehensive analysis of the role of glycosylation and cholesterol on the ability of CDTs from several bacterial species, including A. actinomycetemcomitans. C. jejuni, E. coli, and H. ducreyi, to intoxicate different cell types (15). The authors found that CDT from C. jejuni-induced CHO-K1 cell intoxication was much less efficient than the intoxication of CHO-K1 cells with other CDTs or the intoxication of other cell types with C. jejuni CDT. The authors also demonstrated that C. jejuni CDT-induced cell cycle arrest of CHO-K1 cells was not influenced by cholesterol loading but was enhanced by inhibiting glycosylation. In contrast, in this study, we showed that cholesterol plays a crucial role in the binding of C. jejuni CDT with CHO-K1 cells. This discrepancy may be due to the different concentrations of C. jejuni CDT used in these studies. We applied a higher concentration of C. jejuni CDT (200 nM) than that used in the study of Eshraghi et al. (50 nM), in which cell cycle arrest in CHO-K1 cells became more apparent under the higher concentration of CDT, suggesting that a higher concentration of CDT will advance the understanding of CDT-producing bacteria. Very likely, this outcome will enhance the development of novel therapeutic strategies to prevent or cure diseases caused by these bacterial pathogens.

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We have no conflicts of interest to declare for this work.

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