The role of cholesterol-sphingomyelin membrane nanodomains in the stability of intercellular membrane nanotubes

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Abstract: Intercellular membrane nanotubes (ICNs) are highly curved tubular structures that connect neighboring cells. The stability of these structures depends on the inner cytoskeleton and the cell membrane composition. Yet, due to the difficulty in the extraction of ICNs, the cell membrane composition remains elusive. In the present study, a raft marker, ostreolysin, revealed the enrichment of cholesterol-sphingomyelin membrane nanodomains along ICNs in a T24 (malignant) urothelial cancer cell line. Cholesterol depletion, due to the addition of methyl-β-cyclodextrin, caused the dispersion of cholesterol-sphingomyelin membrane nanodomains and the retraction of ICNs. The depletion of cholesterol also led to cytoskeleton reorganization and to formation of actin stress fibers. Live cell imaging data revealed the possible functional coupling between the change from polygonal to spherical shape, cell separation, and the disconnection of ICNs. The ICN was modeled as an axisymmetric tubular structure, enabling us to investigate the effects of cholesterol content on the ICN curvature. The removal of cholesterol was predicted to reduce the positive spontaneous curvature of the remaining membrane components, increasing their curvature mismatch with the tube curvature. The mechanisms by which the increased curvature mismatch could contribute to the disconnection of ICNs are discussed.

Keywords: cyclodextrins, T24 urothelial cancer cell line, intercellular membrane nanotubes, cholesterol-sphingomyelin membrane nanodomains

Introduction

Intercellular membrane nanotubes (ICNs), as a novel intercellular communication mechanism, are extensively studied both functionally and structurally.¹−⁷ Previously, it has been shown that ICNs are stabilized not only by the inner cytoskeleton but also by the membrane composition.⁸−¹⁰ Yet, due to their fragility, it is impossible to separate the ICNs from the rest of the cell, and therefore their membrane composition remains unknown. To shed light on the ICN membrane composition, it is possible to use membrane raft markers, for the binding of different lipids, proteins, and their complexes.¹¹−¹⁶ In the present study, the addition of a raft marker, ostreolysin (Oly), revealed the presence of cholesterol-sphingomyelin membrane nanodomains throughout the plasma membrane and ICNs. The role of cholesterol-sphingomyelin nanodomains in the stability of ICNs was explored using a cholesterol depletion agent and a cholesterol-free growth medium.

Oly belongs to a novel protein family called aegerolysins (PF06355; IPR 009413), which are known for their cytolytic properties.¹⁷,¹⁸ The binding of Oly evokes the formation of distinctive pores that induce cell lysis.¹⁹ Fluorescent mutants devoid of lytic activity, or fluorescent antibody-labeled proteins at their sublytic concentrations, have been used for the structural investigation of biological membranes.¹¹−¹⁶ In particular,
it has been revealed that Oly interacts specifically with cholesterol-sphingomyelin membrane nanodomains,\textsuperscript{19–21} which do not colocalize with caveolin and cholera toxin B membrane nanodomains.\textsuperscript{11} Oly differs from other raft markers, since it can sense the interaction between two lipids (ie, cholesterol and sphingomyelin), and does not bind to any lipid in its pure form.

While cholesterol-enriched membrane nanodomains have been found to be important for different biological functions, such as exocytosis and endocytosis, signal transduction, pathogen entry, and attachment of various ligands,\textsuperscript{22–24} their contribution to the stability of highly curved membrane regions (eg, ICNs) are not fully understood. A cholesterol molecule has a small hydrophilic headgroup and a relatively short hydrophobic tail, which enables it to fill gaps in highly curved cell membrane regions.\textsuperscript{25,26} The stability of a highly curved ICN might depend on the relative concentration of cholesterol. In mixed membrane models, the addition of cholesterol facilitates the formation of an intermediate phase called liquid ordered phase,\textsuperscript{27,28} exhibiting more rigidity than the surrounding liquid disordered nanodomains.\textsuperscript{29} Therefore, it is possible that the stability of the ICN may depend on the distribution of liquid ordered membrane nanodomains enriched in cholesterol. In the literature, liquid ordered membrane nanodomains were previously denoted as membrane rafts.\textsuperscript{22,23} Membrane rafts are defined as small (10–300 nm) in diameter, heterogeneous, and highly dynamic domains. Upon stimulation, small rafts can be stabilized to form larger platforms through protein–protein and protein–lipid interactions.

Cyclodextrins (CDs) are a family of cyclic compounds made up of oligosaccharides that absorb cholesterol from the cell membrane. The inner core of the CD toroidal structure is much less hydrophilic than the outer surface, which facilitates the absorption of cholesterol from the cell membrane.\textsuperscript{30,31} On the other hand, the release of phospholipids from cell membranes under exposure to CD is only minimal. For example, less than 5% of the initial cell-associated choline-labeled phospholipids was released after 1-hour exposure with 1 mM CD in various Chinese hamster ovary cells and, at most, 2% of cellular phospholipids was released after 5-hour incubation with 5 mM CD from L-cells.\textsuperscript{32,33} Besides their effect on membrane composition, other studies investigated the effect of CDs on cellular communication and cell shaping, demonstrating a decrease in the frequency of endocytosis and in the numbers of membrane invaginations.\textsuperscript{34–36}

Herein, the effects of cholesterol depletion on the mesenchymal shape of a T24 cell and on the density of ICNs are reported. It is revealed that either methyl-\(\beta\)-cyclodextrin (m\(\beta\)CD) treatment or the growth in cholesterol-free medium cause a significant reduction in the density of ICNs. The dynamic morphological alterations are explored using live cell imaging and confocal microscopy. A computational model was constructed to investigate the underlying mechanisms accounting for the instability of tubular structures following the application of a cholesterol depletion agent.

Materials and methods

Urothelial cell culture

Urothelial cell line T24 was cultured in a 1:1 mixture of advanced Gibco\textsuperscript{\textregistered} Dulbecco’s modified Eagle’s medium (Invitrogen Life Technologies, Carlsbad, CA) and Ham’s F-12 (Sigma-Aldrich Corporation, St Louis, MO), supplemented with Gibco 10% fetal bovine serum, 5 \(\mu\)g/mL insulin, 5 \(\mu\)g/mL transferrin, 100 mg/mL hydrocortisone, and 5 ng/mL selinite (Invitrogen), as well as 1800 U/mL cristacyclin (Pliva, Zagreb, Croatia) and 0.222 mg/mL streptomycin sulfate (Fatol Arzneimittel GmbH, Schiffweiler, Germany). Cells were incubated at 37\(^\circ\)C in a humidified incubator (Heracell, Heraeus, Germany) in an atmosphere of 5% carbon dioxide. One day prior to experiment, cells were seeded onto sterile glass coverslips (Brand GmbH, Wertheim, Germany) at approximately 70%–80% confluence and incubated overnight at 37\(^\circ\)C. In some experiments, cells were grown in a medium with lipoprotein poor serum (LPPS medium) purchased from Thermo Scientific HyClone (South Logan, UT).

m\(\beta\)CD

The growth medium of overnight cultures was replaced with cholesterol free media (LPPS) with 1 mM, 2 mM, 4 mM, 6 mM, 8 mM, 10 mM, 12 mM, 16 mM, or 20 mM m\(\beta\)CD (Sigma-Aldrich). Cells were incubated in m\(\beta\)CD for 2 hours or 4 hours, depending on the experiment. Afterwards, incubation medium was removed, cells were fixed in 2% paraformaldehyde (Merck, Hohenbrunn, Germany; 37\(^\circ\)C) for 20 minutes, washed twice in phosphate-buffered saline (PBS; Sigma-Aldrich 37\(^\circ\)C) for 15 minutes, and then analyzed (see below).

Quantification of cellular cholesterol

Lipids from cell homogenates were extracted according to the method of Bligh and Dyer.\textsuperscript{37} Briefly, for each 200 \(\mu\)L of volume, a 1:2 mixture of chloroform:methanol (volume/volume) was added. The chloroform layer was dried under a stream of nitrogen gas and lipids redissolved in isopropyl alcohol. The total cholesterol content was assayed using the
Cholesterol Reagent kit (Thermo Fisher Scientific, Vantaa, Finland) according to the manufacturer’s instructions. In parallel, the protein concentration of 50 μl of each sample was assayed with the Bradford method. The ratio of cholesterol concentration to protein concentration was represented as mg of cholesterol/mg of cell protein.

Oly labeling, actin labeling, antibodies, and microscopy
Oly was isolated from fresh fruiting bodies of *Pleurotus ostreatus* as previously described. The purity of Oly was checked by polyacrylamide gel electrophoresis. The protein concentration was determined spectrophotometrically using the BCA Protein Assay Reagent (Thermo Scientific Pierce Protein Research Products, Rockford, IL). After isolation, the protein was desalted and kept frozen (−20°C) in aliquots in 140 mM sodium chloride, 1 mM ethylenediaminetetraacetic acid, 20 mM tris(hydroxymethyl)aminomethane hydrochloride buffer (pH 8.0). Rabbit anti-Oly primary antibodies were prepared as previously described. T24 cells grown on cover-slips were incubated with 2.5 μg/mL of Oly for 30 minutes at 37°C. After fixation in 4% paraformaldehyde, washing with PBS, and blocking with 2% bovine serum albumin (Sigma-Aldrich) with 0.2% sodium azide, (Fluka Chemie, Buch, Switzerland) primary Oly antirabbit antibodies (1:2500) and then Molecular Probes® Alexa Fluor® 555-conjugated secondary antibodies (1:1000) (Invitrogen) were added. Coverslips were washed and mounted in VECTASHIELD® with 4′,6-diamidino-2-phenylindole (Vector Laboratories, Burlingame, CA).

Actin labeling was performed in 16.7 μg/mL phalloidin (phalloidin-fluorescein isothiocyanate) (Sigma-Aldrich) in 20% methanol (Carlo Erba Reagents, Milan, Italy) in PBS for 30 minutes. For preparations which were labeled for Oly, actin labeling with phalloidin-tetramethylrhodamine isothiocyanate (Sigma-Aldrich) was performed after goat antirabbit secondary antibody incubation and 10 minutes washing in PBS. Afterwards, coverslips were decanted and embedded in VECTASHIELD-4′,6-diamidino-2-phenylindole and analyzed in a fluorescence microscope (Axio™ Imager Z1; Carl Zeiss AG, Oberkochen, Germany).

Phase-contrast and fluorescence image acquisition
Cells were analyzed in a fluorescence microscope (Axio Imager Z1). Phase-contrast images were taken with 63×-water objective (numerical aperture 0.95) and fluorescence images with Plan Apochromat® (63×, oil/numerical aperture 1.4; Carl Zeiss).

Morphometric analysis
Sampling was performed as follows: cover glass with fixed cells was imaged with water immersion objective (63×). Images were taken at every second visual field, second horizontal, and second vertical axis. Afterwards, 20 randomly chosen images were analyzed for each treatment by counting the cells and ICNs per image. Only membrane nanotubes that made contact between neighboring cells were counted as ICNs. The ICN density, ie, the number of ICNs per cell, was calculated as follows: ICN density = (total number of ICNs/total number of cells). Data are reported as mean ± standard deviation in tables and text.

Live cell imaging
T24 cells were cultured on glass bottom dishes (MatTek Corporation, Ashland, MA). Time-lapse imaging was obtained on an LSM 510 (Carl Zeiss) confocal microscope using transmission light (oil objective 63×). Initially, images of control were taken (data not shown), then the medium was exchanged with cholesterol-free growth medium treated with 5 mM mβCD and time sequences collected every 18 seconds for 2 hours.

Results
Oly and other raft markers
Previously, it has been demonstrated that Oly preferentially binds to cholesterol-sphingomyelin membrane nanodomains. Figure 1B shows the binding of Oly to cholesterol-sphingomyelin membrane nanodomains throughout the plasma membrane and ICNs in T24 cells. Oly preferentially binds rafts enriched with cholesterol and sphingomyelin molecules. Oly was strongly labeled in some of the T24 cells (Figure 1B), which could be due to differences in the concentration of cholesterol-sphingomyelin rafts. Along the ICNs, there was no binding of caveolin-1 and flotillin-1 raft markers, and very little binding of ganglioside GM1 raft marker (data not shown).

The depletion of cholesterol using mβCD and LPPS revealed reduced binding of Oly to the cell surface (Figure 1D), which demonstrated the disappearance of cholesterol-sphingomyelin membrane nanodomains that Oly binds to in the cell membrane. Finally, due to the relatively low concentration of mβCD (1 mM) and the short treatment time (2 hours), there was neither an effect on the cell morphology nor on the ICN density (Figure 1C and D).
mβCD treatment

CD treatment has been shown to drive the transfer of cholesterol from the outer membrane leaflet of the cell and from fluid areas to mβCD molecules. In the present study, the depletion of cholesterol from the plasma membrane was verified using the Bligh and Dyer method. It was revealed that the treatment by mβCD reduced the concentration of cholesterol in cell membranes. The relative concentration of cholesterol (mg cholesterol/mg protein) was 0.006 in the control experiment, whereas following mβCD treatment (4 mM for 2 hours), the relative concentration fell to 0.002.

Figure 2 shows phase-contrast images of T24 cells grown in LPPS for 2 hours and treated with increasing concentrations of mβCD (from 2 mM to 20 mM). Following treatment, cells were more separated, and the extent of cell separation was a function of mβCD concentration. Furthermore, it was revealed that 2 mM mβCD did not have a significant effect on the ICN frequency (Figure 2C). However, above a concentration of 4 mM mβCD, the retraction of ICNs became considerable, and the cell edge surface became smoother, demonstrating the retraction of short and long membrane protrusions (Figure 2D–J). These results revealed that the depletion of more cholesterol led to more significant morphologic changes.

Actin and DNA labeling

The formation of actin stress fibers might be important for cell reshaping and cell contraction. Following cell growth in LPPS for 4 hours, actin labeling of T24 cells demonstrated the smoothing of the cell surface as well as the formation of actin stress fibers (Figure 3C). The formation of actin stress fibers might facilitate static cell contraction, which could reduce the cell size and smooth out the cell surface. Similar morphologic changes were observed upon 4 mM mβCD treatment (Figure 3D and E). By adding a DNA label (4′,6-diamidino-2-phenylindole), it was observed that the integrity of the cell nuclei, after mβCD treatment, remained intact (Figure 3F–J).

Morphometric analysis

To determine the statistical effect of mβCD treatment in T24 cells, the number of cells and the number of ICNs were counted in acquired images. The ICN density was calculated in each image. Summary statistics for the differences in ICN density upon mβCD treatments of different concentrations for 2 hours are shown in Figure 4. It was revealed that the ICN density was significantly (P < 0.05) reduced following the application of 4 mM mβCD, and that the average number of ICNs fell dramatically upon an increase in the concentration.

Figure 1 Ostreolysin marks the presence of cholesterol-sphingomyelin membrane nanodomains along intercellular membrane nanotubes in T24 cells. The labeling of actin filaments highlights the location of intercellular membrane nanotubes (see close up) between T24 (malignant) cells of urothelial origin grown in lipoprotein poor serum medium (A). The addition of ostreolysin reveals specific binding to cholesterol-sphingomyelin membrane nanodomains along the cell surface and along the intercellular membrane nanotubes (B). The growth of T24 cells in lipoprotein poor serum medium treated with 1 mM methyl-β-cyclodextrin for 2-hour causes the formation of actin stress fibers (C) as well as the dispersion of cholesterol-sphingomyelin membrane nanodomains (D). Note that the intercellular membrane nanotubes are still present following the application of methyl-β-cyclodextrin in low concentration (1 mM) (C), and that ostreolysin labeling on the cell surface is much reduced (D).

Abbreviations: LPPS, lipoprotein poor serum; mβCD, methyl-β-cyclodextrin; Oly, ostreolysin.
Figure 2 Phase-contrast images of T24 cells revealing the loss of intercellular membrane nanotubes upon increasing concentration of methyl-β-cyclodextrin treatment. The images show the growth of T24 cells in control (A) and cholesterol-free medium (lipoprotein poor serum) (B), and the morphological changes of T24 cells in lipoprotein poor serum following 2–20 mM methyl-β-cyclodextrin 2-hour treatment (C–J). The intercellular membrane nanotubes, observed following 4 mM methyl-β-cyclodextrin treatment, are thinner, longer (C; see arrow), and less frequent than in the control experiment (A; see arrow). The removal of cholesterol from the membrane causes the inward contraction of the cell plasma membrane (D and E). The formation of large vesicular dilations along the intercellular membrane nanotube (E; see arrow) could be due to the curvature mismatch between the remaining membrane components and the tube curvature. Above 6 mM methyl-β-cyclodextrin, the cells become isolated and spherical (F–J).

Abbreviation: mβCD, methyl-β-cyclodextrin.
of mβCD from 4 mM to 6 mM (Figure 4). This shows that there might be a threshold of cholesterol content in the cell membrane, below which there is a dramatic decline in ICN density.

The coupling between ICN density and cell shaping

To shed light on the dynamic changes in cell morphology following cholesterol depletion, live cell imaging techniques were employed using confocal laser scan microscopy. The hypothesis was that the disruption of ICNs and the change from a polygonal to a spherical cell shape were mechanically coupled. Using a confocal microscope (LSM 510) and time-lapse imaging, the cell morphologic changes were tracked during treatment of 5 mM mβCD. Figure 5A shows that ICNs are disrupted somewhere in the middle, and then retracted towards the parent cell membrane. Furthermore, the disconnection of ICNs caused the abrupt cell separation and the rounding of a T24 cell (Figure 5).

**Figure 3** Fluorescence microscope images of T24 cells revealing the integrity of actin filaments and cell nucleus following the growth in cholesterol-free (lipoprotein poor serum) or methyl-β-cyclodextrin treatments. The actin labeling of T24 cells in the control experiment reveals the cell surface ruffling (A). The growth of T24 cells in lipoprotein poor serum causes the smoothing of the cell surface (retraction of the membrane protrusions) as well as the formation of actin stress fibers, which is observed more after 4-hour (C) than 2-hour (B) treatment. The morphological changes in T24 cells following methyl-β-cyclodextrin treatment are similar to the changes observed following growth in lipoprotein poor serum (D and E). Using a DNA label (4’,6-diamidino-2-phenylindole), the integrity of the cell nuclei in the corresponding lipoprotein poor serum and methyl-β-cyclodextrin treatments is revealed (F-J).

**Abbreviations:** DAPI, 4’,6-diamidino-2-phenylindole; h, hours; LPPS, lipoprotein poor serum; mβCD, methyl-β-cyclodextrin.

**Figure 4** Summary statistics of changes in intercellular membrane nanotube density among T24 cells grown in different growth mediums and following different methyl-β-cyclodextrin concentration treatments. The statistical differences are calculated with respect to the control case. Note that the reduction in intercellular membrane nanotube density becomes significant from 4 mM methyl-β-cyclodextrin 2-hour treatment and highly significant above this concentration.

**Note:** Data are mean ± standard deviation; *P < 0.05; **P < 0.01.

**Abbreviations:** ICN, intercellular membrane nanotube; mβCD, methyl-β-cyclodextrin.
Possible mechanisms for mβCD-induced morphological changes of T24 cells

There are three possible mechanisms responsible for the observed reduction in the ICN density during mβCD treatment. The first mechanism is that the “gaps” created in the cell membrane, due to cholesterol depletion from the external leaflet and fluid areas, were filled by the surrounding membrane, thereby causing the contraction of the cell membrane and the partial retraction of membrane protrusions. The second possible mechanism is based on the positive spontaneous curvature of a cholesterol-phospholipid complex (nanodomain) (Figure 6). This spontaneous curvature could be due to an area difference between the two membrane leaflets, assuming that more cholesterol molecules might reside in the outer than in the inner membrane leaflet (Figure 6A), or due to a spontaneous curvature of the cholesterol-phospholipid complex (Figure 6B). The plasma membrane of the...
human erythrocyte contains in the outer membrane leaflet 65%–75% of the phosphatidylcholine and >85% of the sphingomyelin, while in the inner membrane leaflet there is 80%–85% of the phosphatidylethanolamine and >96% of the phosphatidylserine. Since the outer membrane leaflet is enriched in sphingomyelin, the preferred interaction of cholesterol with sphingomyelin may justify the assumption of cholesterol residing more in the outer than in the inner membrane leaflet of the ICN. The third possible (complementary) mechanism is that cell separation stretches the ICNs, until the ICNs are disrupted in the middle or detached from the parent cell membrane.

To minimize the bending energy costs during the formation of a thin tubular structure, cholesterol molecules could accumulate in the outer membrane leaflet. Therefore, the depletion of cholesterol from the outer membrane leaflet would increase the mismatch between the tube curvature and the spontaneous curvature of the remaining membrane nanodomains.

The model of ICN stability and theoretical predictions

Phase-contrast images of T24 cells grown in LPPS and mβCD for 2 hours revealed that during cell separation the ICNs became thinner and longer and, eventually, disrupted (Figures 2 and 5). The aim of the present model was to elucidate the possible underlying mechanisms responsible for the disruption of ICNs. The computational model was an extension of a previous model. The model included only the ICN isolated, ie, it excluded the membrane of the parent cells (Figure 7C). The ICN was modeled as an axisymmetric shape of cylindrical symmetry. For the sake of simplicity, the membrane was composed of only one component (nanodomain). The component was a cholesterol- and sphingomyelin membrane nanodomain, consisting of cholesterol, sphingomyelin, protein, and other lipid molecules (Figure 7A). The spontaneous curvature of the nanodomain was assumed to be positive, which was based on either nonzero area difference between the two
membrane leaflets (Figure 6A) or the intrinsic curvature of the cholesterol-phospholipid complex (Figure 6B). The underlying assumption of the model was the asymmetry in the cholesterol content between the inner and outer membrane leaflets along the axisymmetric tubular structure, such that there were more cholesterol molecules in the outer than in the inner membrane leaflet of the ICN (Figure 6), influencing the nanodomain intrinsic (spontaneous) curvature. The sign of the membrane curvature was defined with respect to the outer membrane leaflet. The membrane bilayer was modeled as a single entity.

The membrane free (bending) energy is considered within continuum approach in the form:  

$$ E = \int \left( \kappa (H - H_m(\phi))^2 + \kappa (D - D_m(\phi))^2 + \sigma \right) dA $$  

(1)

where $\kappa$ is the membrane bending rigidity, $\phi$ is the relative density of cholesterol molecules in the outer membrane leaflet, $H$ is the mean membrane curvature, $D$ is the local membrane curvature deviator, $H_m(\phi)$ is the intrinsic (spontaneous) mean curvature of the nanodomain, $D_m(\phi)$ is the intrinsic (spontaneous) membrane curvature deviator of the nanodomain, $\sigma$ is the Lagrange multiplier, and $dA$ is the infinitesimal membrane area element. The integral is taken over the surface of the axisymmetric tube. The mean membrane curvature takes into account the two principal curvatures of the axisymmetric cylindrical structure. The difference between the membrane curvature and the membrane intrinsic (spontaneous) curvature determines the energy cost for bending the membrane away from its favorable curvature.  

The intrinsic (spontaneous) curvatures of the membrane nanodomain depend on the relative density of cholesterol molecules in the outer lipid layer, as follows:

$$ H_m(\phi) = H_{m,0} + \frac{H_{m,c}}{H_{m},0} \phi $$

$$ D_m(\phi) = D_{m,0} + \frac{D_{m,c}}{D_{m,0}} \phi $$  

(2)

where $H_{m,c}$ and $D_{m,c}$ are the intrinsic (spontaneous) mean and deviatoric curvature of the cholesterol-free nanodomain, respectively, and $H_{m,0}$ and $D_{m,0}$ are the intrinsic (spontaneous) mean and deviatoric curvature of the cholesterol-replenished (maximum concentration) nanodomain, respectively.

The equation of motion of the membrane height deflection along the cylinder main symmetry axis is given by:

$$ \frac{\partial h(z)}{\partial t} = - \frac{\partial E}{\partial h(z)}, $$  

(3)

where $\phi$ is the friction coefficient describing the drag of the fluid surrounding the membrane, $t$ is time, and $E$ is redefined as the membrane free energy per unit area. The system is investigated in the limit of small membrane deformation from the uniform radius of the ICN. By linearization of Equation (3), the following equation was obtained:

$$ \frac{\delta h(z)}{\delta t} = \int (U + L(h)\delta + O(\delta^2))dA, $$  

(4)

where the function $L(h)$ describes the force in the limit of small and linear membrane undulations. The integral sign is for the calculation of the average force density. The force acting on the membrane in the equilibrium state is described by $U$. The equilibrium radius $R_{eq}$ is obtained by assuming that the undulation $U$ equals zero, which yields:

$$ R_{eq} = \frac{\sqrt{\kappa}}{\sqrt{2\sigma + (H_m(\phi)^2 + D_m(\phi)^2)\kappa^2}} $$  

(5)

It is important to note that the axisymmetric model of the ICN is stable even without anisotropy, since it is not considered as a membrane protrusion.

An aim of this study was to analyze the effects of changes in the relative density of cholesterol on the equilibrium radius of the tube. Figure 7D shows the equilibrium radius as a function of the relative density of cholesterol. The equilibrium radius remains low at 0.15 $\mu$m for a relative density of 0.3. When the relative density is reduced below 0.2, the equilibrium radius is increased rapidly. A plausible range of cholesterol content in the plasma membrane is between 20%–30%. According to calculations, the cholesterol content might be a critical parameter, having a role not only in the stability of membrane nanodomains but also affecting the radius of the highly curved structure. The complete depletion of cholesterol ($\phi = 0$) from the outer leaflet of the tubular membrane is predicted to considerably increase the curvature mismatch between the spontaneous curvature of the remaining membrane components and the tube curvature. To reduce the curvature mismatch, vesicular dilations could be formed during the retraction of the ICN (Figure 2E).

**Discussion**

This study demonstrates the presence of cholesterol-sphingomyelin membrane nanodomains along ICNs, which was revealed by the addition of membrane raft marker Oly (Figure 1). However, the ICNs were negative for the binding of other membrane raft makers such as flotilin-1, caveolin-1,
and ganglioside GM1. The removal of cholesterol from the cell membrane by the application of mβCD led to the smoothing of the cell surface, the separation of cells, the stretching of ICNs, and the retraction of disconnected ICNs (Figure 2). Following cholesterol depletion, the labeling of actin filaments revealed the formation of actin stress fibers and the reorganization of the actin cytoskeleton (Figure 3). Statistical analysis revealed significant \( P < 0.01 \) reductions in the density of ICNs upon exposure to high concentrations of mβCD (Figure 4). The change of cell shape from polygonal to spherical would increase the distance between neighboring cells, stretching the connected ICNs. A computational model was constructed to investigate the contributions of changes in the spontaneous (intrinsic) curvature of cholesterol-sphingomyelin membrane nanodomains to the shape and stability of ICNs (Figures 6 and 7). It was theoretically predicted that the depletion of cholesterol from the outer membrane leaflet would favor a larger radius of the tubular structure. Actually, the reduction in the ICN radius was observed (Figure 5), which increases the curvature mismatch between the tube curvature and the spontaneous curvatures \( (H_m(\phi) \text{ and } D_m(\phi)) \) of the remaining membrane components. This curvature mismatch could be reduced by the formation of large vesicular dilations along the ICN (Figure 2E), and by the detachment of ICNs from the parent plasma membranes (Figure 5A).

CDs have been mainly used for the rapid extraction of cholesterol molecules from cell membranes,\(^{39,40}\) while the release of phospholipids was only minimal.\(^{32,33}\) Previous studies have shown that the depletion of cholesterol by CDs might lead to a reduction in endocytosis and membrane invagination.\(^{34–36}\) Yet, the underlying mechanism is probably different from the one in the present model, since membrane invaginations are enriched in cholesterol-based caveolin nanodomains, which do not contain sphingomyelin molecules. The underlying assumption of the present model is that the cholesterol-sphingomyelin complex has a positive spontaneous curvature, whereas, cholesterol-based caveolin nanodomains may have a negative spontaneous curvature.

The depletion of cholesterol from the outer membrane leaflet may reduce the area difference between the two membrane leaflets. As a result, the plasma membrane would favor more planar-like conformations, explaining the smoothing of the cell surface and the retraction of disconnected ICNs (Figure 2 and Figure 5A). Since it has been recently shown that cholesterol-sphingomyelin nanodomains were essential for microtubule-based protrusion growth,\(^{44}\) it could be also possible that the stability of the ICN depends on the binding of microtubules to the ICN membrane through cholesterol-based membrane nanodomains.

According to the theory of isotropic membrane elasticity,\(^{35,46}\) the cell tubular membrane protrusion cannot be stabilized by isotropic membrane components alone,\(^{33}\) but requires the accumulation of anisotropic membrane nanodomains.\(^{5,8,10,47,48}\) Previous theoretical studies provide insight into the stability of the tubular membrane protrusion following the disassembly of the inner rod cytoskeleton,\(^{4}\) or into the stability of actin-free filopodia.\(^{10,49}\) Accordingly, it is possible that the accumulation of cholesterol-sphingomyelin membrane nanodomains of anisotropic spontaneous curvature may contribute to the tube stability by overcoming the decrease in configurational entropy during the process of lateral sorting of the nanodomains.\(^{8,6,10,48–50}\) Therefore, the loss of cholesterol-sphingomyelin rafts that may be connected to reduced intrinsic anisotropy of the tube membrane might cause dynamic instability leading to the retraction of disconnected ICNs.

In a previous study, it has been shown that cholesterol-sphingomyelin membrane domains protrude out of the membrane surface,\(^{51}\) thereby increasing the lateral tension between the protruding ordered nanodomain and the neighboring disordered nanodomains. The increased membrane tension can be reduced by cholesterol molecules bridging the membrane height difference between the ordered and disordered nanodomains. It is also possible that the high curvature along ICNs would introduce additional membrane tension. Consequently, the cholesterol content in cholesterol-sphingomyelin membrane nanodomains along ICNs could be greater than in similar nanodomains on the cell (spherical) plasma membrane.

In conclusion, the present experimental and theoretical results suggest that the transfer of cholesterol from the outer membrane leaflet of cell membranes of malignant T24 cells to CDs molecules causes the smoothing of the cell surface accompanied by the retraction of ICNs. A future study will investigate whether these morphologic alterations are due to the amounts of membrane lipids being depleted or due to the change in the spontaneous curvature of the cholesterol-free membrane regions. The possibility of different membrane nanodomain composition organized by the high cylindrical curvature of ICNs could be investigated using cholesterol/sphingomyelin probe Oly.

As the ICNs are supposed to represent an important communication system enabling metastatic cancer cells to explore their surroundings,\(^{7}\) their reduction by cholesterol removal might represent a new target for anticancer therapy.
The exploitation of ICNs for the movement of pathogens (e.g., viral particles such as human immunodeficiency virus) on top of the membrane nanotubes from infected to noninfected cells might be dependent on cholesterol-based membrane nanodomains. The contributions of hopanoids (the bacterial equivalent of cholesterol) to the stability of membrane nanotubes that confer antibiotic resistance could be explored in a future study. The present results regarding the functional link between the membrane composition and the stability of the ICN may shed light on the intercellular communication between neighboring cells.

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Disclosure
The authors report no conflicts of interest in this work.

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