

A scissors mechanism for stimulation of SNARE-mediated lipid mixing by cholesterol

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Neurotransmitter release at the synapse requires membrane fusion. The SNARE complex, composed of the plasma membrane t-SNAREs syntaxin 1A and SNAP-25 and the vesicle v-SNARE synaptobrevin, mediates the fusion of 2 membranes. Synaptic vesicles contain unusually high cholesterol, but the exact role of cholesterol in fusion is not known. In this study, cholesterol was found to stimulate SNARE-mediated lipid mixing of proteoliposomes by a factor of 5 at a physiological concentration. Surprisingly, however, the stimulatory effect was more pronounced when cholesterol was on the v-SNARE side than when it was on the t-SNARE side. Site-directed spin labeling and both continuous wave (CW) and pulsed EPR revealed that cholesterol induces a conformational change of the v-SNARE transmembrane domain (TMD) from an open scissors-like dimer to a parallel dimer. When the TMD was forced to form a parallel dimer by the disulfide bond, the rate was stimulated 2.3-fold even without cholesterol, supporting the relevance of the open-to-closed conformational change to the fusion activity. The open scissors-like conformation may be unfavorable for fusion and cholesterol may relieve this inhibitory factor.

EPR | membrane fusion | syntaxin | synaptobrevin

In the neuron, synaptic vesicles fuse with plasma membrane to release neurotransmitters into the synaptic cleft (1–4). Membrane fusion involves extensive bilayer remodeling, which requires free energy or an effective catalyst (5–6). SNAREs (soluble *N*-ethylmaleimide-sensitive factor attachment protein receptors) are believed to be the fusion machine that serves as the energy source (7–8) or catalyst that lowers the fusion energy barrier (9). At the onset of fusion, vesicle associated (v-) SNARE synaptobrevin (or VAMP2) combines with target membrane (t-) SNAREs Syntaxin 1A and SNAP-25 to form a helical bundle that bridges 2 membranes, facilitating fusion (10–14).

Cholesterol is a major component of the plasma and the vesicle membranes, composing as much as 30–40 mol % of the total lipids (15–16). Cholesterol is structurally rigid and known to harden the membrane significantly, making it resistant to being bent or deformed. However, there is evidence that cholesterol accelerates membrane fusion that involves a series of bilayer deformations (17–20). The physical origin of cholesterol's positive role can be found partly in its spontaneous negative curvature such that it prefers to be in the negatively curved (concave) surface but dislikes being in the positively curved (convex) surface (21).

Despite progress in understanding the effects of cholesterol on the bilayer curvature, not much is known about its influence on the lateral distribution of SNAREs in the membrane or on structures of their transmembrane domains (TMDs). A series of recent reports addresses the importance of the clustering of SNAREs at the fusion site for successful fusion (19, 22–24). In addition, there is evidence that the SNARE TMDs are essential for formation of the fusion pore (25–27). Considering cholesterol's structural rigidity and its nonideal mixing with phospholipids (28), we speculate that both the lateral distribution and the conformation of the SNARE TMDs are affected significantly by the presence of cholesterol. Indeed, it has been shown that cholesterol is a key component to induce lateral clustering of Syntaxin 1A (19).

In this work, we attempt to correlate the conformational changes of the SNARE TMDs induced by cholesterol to the fusion activity of neuronal SNAREs. The fluorescence lipid-mixing assay was used to investigate the effect of cholesterol on SNARE-mediated membrane fusion. The results show that lipid mixing was promoted significantly when cholesterol is on the v-SNARE-carrying vesicles but not as much when it is on t-SNARE-carrying vesicles. Using site-directed spin labeling (SDSL) and both continuous wave (CW) and pulsed-EPR spectroscopy, we found a significant conformational change of the TMD dimers of v-SNARE synaptobrevin (or VAMP2) that is induced by cholesterol. This change resembles a scissors-type motion of the TMD dimers. The relevance of this structural change to the fusion activity was verified by using the disulfide cross-linking method.

Results

Physiological Cholesterol Stimulates Lipid Mixing. To investigate the effect of cholesterol on SNARE-mediated lipid mixing, we incorporated appropriate amounts of cholesterol into the POPC/DOPS vesicles to achieve 3 different cholesterol concentrations of 0, 20, and 40 mol %. We kept the negatively charged DOPS level at 15 mol % for all 3 samples. The t-SNARE complex was preformed by mixing Syntaxin 1A and SNAP-25 in the 1:1 molar ratio. The t-SNARE complex was then reconstituted into the vesicles (t-SNARE vesicles). The v-SNARE protein VAMP2 was incorporated into a separate population of vesicles that contained 1.5 mol % each of NBD-PS (1,2-dioleoyl-*sn*-glycero-3-phosphoserine-*N*-(7-nitro-2-1,3-benzoxadiazol-4-yl) and rhodamine-PE (1,2-dioleoyl-*sn*-glycero-3-phosphoethanolamine-*N*-(lissamine rhodamine B sulfonyle) for the fluorescence detection of lipid mixing (v-SNARE vesicles). The lipid-to-protein ratio was kept at 200:1 for all samples (except as noted). To follow the kinetics of lipid mixing, we mixed t-SNARE vesicles and v-SNARE vesicles at 37 °C in the 9:1 ratio. When SNAP-25 was not included, no lipid mixing was detected, as expected (black trace in Fig. 1A). However, significant lipid mixing was observed when SNAP-25 was included in the t-SNARE vesicles (Fig. 1A).

When physiological 40 mol % of cholesterol was included in both vesicles, we observed a 5-fold increase of the initial rate of lipid mixing with respect to that in the absence of cholesterol (Fig. 1B). Interestingly, there was only minor stimulation of lipid mixing when 20 mol % of cholesterol was incorporated in both membranes. Therefore, the results show that the physiological concentration of cholesterol is necessary to produce significant stimulation of lipid mixing.

SNARE-mediated fusion transits through the hemifusion state in which the outer leaflets are merged but the inner leaflets remain

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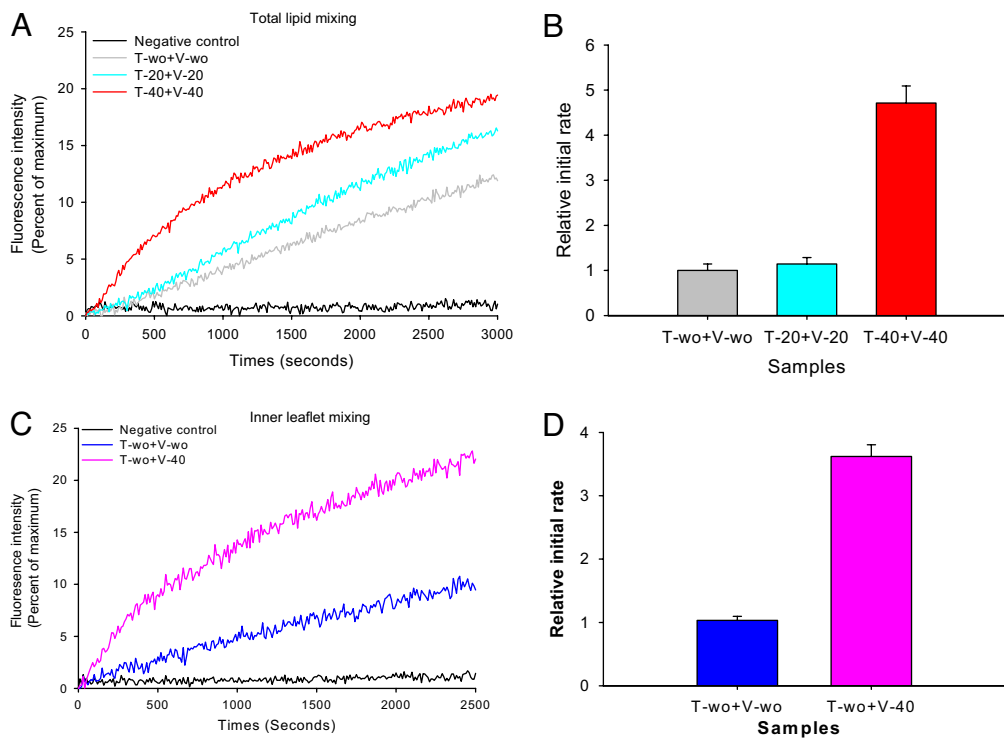


Fig. 1. Lipid-mixing assays at several different cholesterol concentrations. (A) The changes of NBD fluorescence reflecting lipid mixing. The black trace is the control run with the t-liposomes reconstituted with only Syntaxin 1A (without SNAP-25). The gray trace shows lipid mixing for both v- and t-liposomes without cholesterol. The cyan trace is the lipid-mixing assay with 20 mol % cholesterol on both sides. The red trace represents the lipid-mixing assay with 40 mol % cholesterol on both liposomes. (B) The relative initial rates of lipid mixing. The initial rates were determined by fitting the rise of the fluorescence intensity with the linear line near $t = 0$. (C) The inner leaflet mixing assay. The gray trace is the negative control without SNAP25. The blue trace is the fluorescence change representing inner leaflet mixing for the liposomes without cholesterol. The pink trace represents inner leaflet mixing for the liposomes with 40 mol % cholesterol. (D) The relative initial rates for inner leaflet mixing. The error bars were obtained from at least $n = 4$ independent measurements from 4 different preparations.

intact (29–31). To further dissect the effect of cholesterol on hemifusion (outer leaflet mixing) and full fusion (inner leaflet mixing), we performed the inner leaflet lipid-mixing assay (Fig. 1C). We used the modified Meer's method (29) that uses dithionite to selectively reduce the fluorescence donor NBD on only the outer leaflet, which allows the selective detection of inner leaflet mixing. The results show that the rate of inner leaflet mixing was enhanced by a factor of 4 at 40 mol % cholesterol (Fig. 1D), indicating that cholesterol accelerates both hemifusion and full fusion similarly.

Cholesterol Is More Effective When It Is on the v-SNARE Vesicle. A lipidomic study by Jahn and coworkers (16) revealed a surprising result that cholesterol is highly enriched at the synaptic vesicles, constituting ≈ 40 mol % of the total lipids, which is higher than the cholesterol content of the plasma membrane. Thus, we wonder whether the high cholesterol content in the vesicle has any functional implications on SNARE-mediated lipid mixing.

To address this question, we analyzed lipid mixing between the cholesterol-containing v-SNARE vesicles and the cholesterol-absent t-SNARE vesicles and vice versa (Fig. 2A). When 40 mol % of cholesterol was included in the v-SNARE vesicles with no cholesterol on the t-SNARE vesicles, the rate of lipid mixing was $\approx 70\%$ of that for 40 mol % cholesterol on both vesicles (Fig. 2B).

In contrast, when 40 mol % of cholesterol was loaded on the t-SNARE vesicle with no cholesterol on the v-SNARE vesicle, we observed stimulation of lipid mixing by a factor of only 1.7 with respect to the control without cholesterol on both sides. Thus, the result shows that cholesterol on the t-SNARE vesicle is less effective than it is on the v-SNARE side. Such asymmetric effects of cholesterol on SNARE-dependent lipid mixing may not be fully accounted for by the cholesterol's membrane curvature-modifying role. The results may suggest that its impact on the protein structure, particularly on the structure of v-SNAREs, also has important functional implications.

SDSL EPR Reveals the v-SNARE TMD Forms an Open Scissors-Like Dimer. We used SDSL (32–34) and EPR spectroscopy to investigate the conformational changes of the v-SNARE TMD induced by cholesterol. In SDSL, native residues were replaced 1 at a time with cysteines to provide specific labeling sites for the thiol-specific nitroxide spin labels such as methanethiosulfonate spin label (MTSSL) (35, 36). The EPR line shape was sensitive to the motion of the nitroxide as well as to the spin–spin interaction with the neighboring nitroxides. For example, the motional restriction caused by the tertiary interaction with the neighboring helix would broaden the EPR lines. Yet, if the pairs of nitroxides are within 25

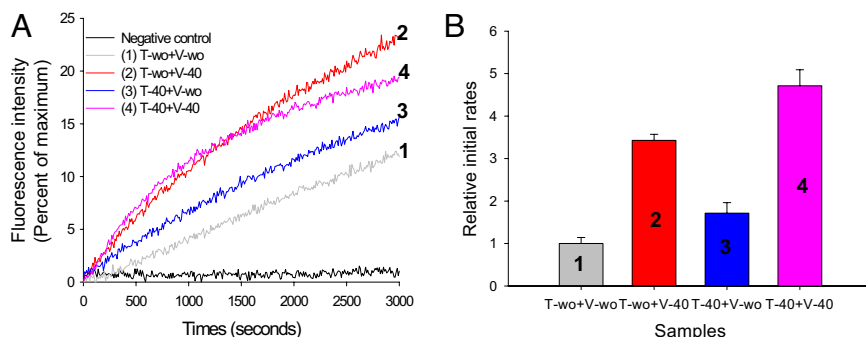


Fig. 2. Lipid-mixing assays with asymmetric distribution of cholesterol. (A) The changes in the NBD fluorescence intensity reflecting lipid mixing. The black trace is the control run with the t-SNARE liposome reconstituted with only Syntaxin 1A (without SNAP-25). The gray trace is for the case without cholesterol. The red trace is for v-liposome with 40 mol % cholesterol but no cholesterol on t-liposome. The blue trace is vice versa. The pink trace is for 40 mol % cholesterol on both sides. (B) The relative initial rates of lipid mixing. The initial rate without cholesterol on both sides was set as 1. The error bars were obtained from at least $n = 4$ independent measurements from 4 different preparations.

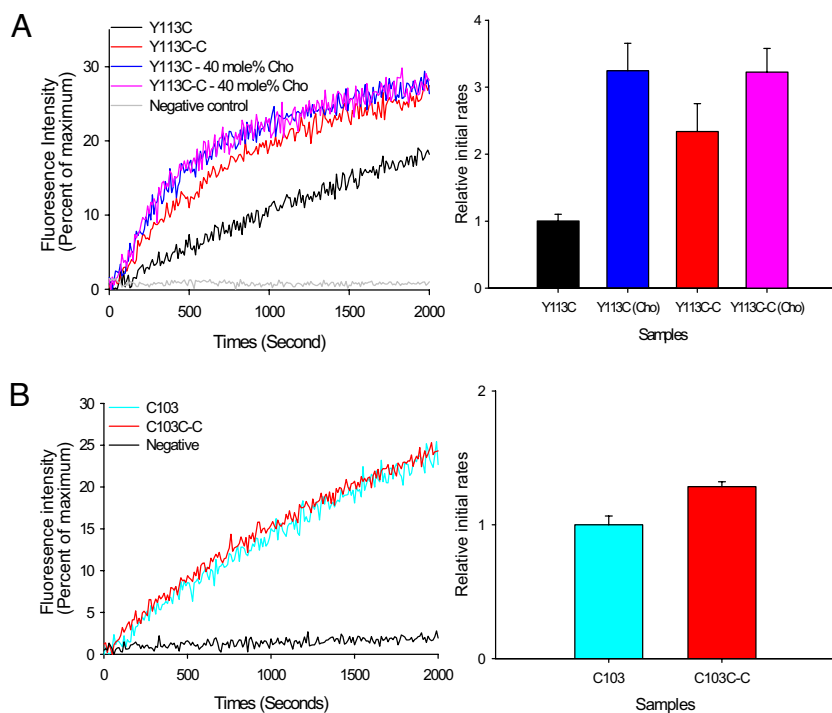


Fig. 4. Lipid-mixing assays for cross-linked VAMP2 wild-type C103 or mutant Y113C. (A) The NBD fluorescence change for cross-linked Y113C-C. (Left) The gray trace is a negative control without SNAP-25. The black trace is for non-cross-linked VAMP2 mutant Y113C without cholesterol. The red trace is for the cross-linked VAMP2 mutant Y113C-C without cholesterol. The blue and pink traces are for the non-cross-linked Y113C and cross-linked Y113C-C in the liposomes with 40 mol % cholesterol, respectively. (Right) The relative initial rates. (B) The NBD fluorescence change for cross-linked C103C-C. (Left) The cyan trace is for the lipid-mixing assay for non-cross-linked VAMP2 C103. The red trace is the cross-linked VAMP2 C103C-C. (Right) The relative initial rates.

whereas the C-terminal halves were splayed apart, resembling an open scissors (Fig. 5A). However, we note that the helices appeared to be loosely held together and so dynamic that the tertiary interaction could not be picked up in the EPR line shape and the distance distributions were broad (SI Text).

Cholesterol Brings the C Termini of the TMDs Closer. When cholesterol was incorporated in the vesicles, we observed small but clear and unambiguous line broadening at several selected positions including residues 99, 102, 106, 113, and 114 (Fig. 3A and D, red traces). This was most likely due to the tertiary contacts between the TMDs. Had the line broadening been simply due to the cholesterol-induced decrease of the membrane viscosity, it would have occurred at all positions and not only to a select few positions. In the low-temperature spectra, the line broadening due to the spin-spin interaction became pronounced at several positions (Fig. 3B, red traces). What is particularly interesting was the conspicuous decrease of distances at the C-terminal positions in the presence of cholesterol (Fig. 3C). The distance profile throughout the TMD was quite consistent with the nearly parallel dimer, with positions 96, 99, 103, 106, 110, and 113 at the dimer interface (Fig. 3D). The TMD dimer in the presence of 40 mol % cholesterol appeared to resemble the closed scissors (Fig. 5B), as opposed to the open scissors-like dimer in the absence of cholesterol.

Is the Cholesterol-Induced Change in the TMD Relevant to the Fusion Activity? The EPR analysis of spin-labeled VAMP2 showed that cholesterol induces an open-to-closed conformational change in the TMD. To verify whether this conformational change is functionally relevant, we used cysteine-cysteine cross-linking. Cysteine-cysteine cross-linking at a C-terminal position such as Y113C, which is a predicted interfacial position, would bring the C-terminal parts of the dimer into contact, forcing the dimer to take the closed form even in the absence of cholesterol. If the open-to-closed conformational change is functionally important, we should observe the stimulation of lipid mixing with the disulfide-linked dimer in the absence of cholesterol. For the mutant Y113C, 40 mol % cholesterol stimulated the initial rate of lipid mixing by a factor of 3, somewhat less than that for wild-type VAMP2.

When VAMP2 Y113C was cross-linked by Cu(II) phenanthroline (Fig. S3) and subjected to the lipid-mixing assay with the vesicles without cholesterol, we observed the increase of the initial rate by a factor of 2.3 (Fig. 4A), generally supporting our prediction. But the stimulation by cross-linking was less than a factor of 3 and was somewhat short of fully accounting for the effect of cholesterol on lipid mixing. Interestingly, however, when cholesterol (40 mol %) was added in the lipid mixing assay with cross-linked Y113C-C, the further gain was only 25% (Fig. 4A), suggesting that the cholesterol-induced closure of the VAMP TMD dimer accounts for approximately three-fourths of the cholesterol effect on SNARE-dependent lipid mixing. The remaining one-fourth may be attributed to other factors such as the cholesterol's spontaneous curvature. Thus, the analysis suggests that the open-to-closed conformational change in the VAMP2 TMD by cholesterol is the major factor that contributes to the stimulation of lipid mixing.

Yet, when N-terminal position C103 was cross-linked, there was little change in the initial rate when compared with that for the non-cross-linked C103 (Fig. 4B). This was again expected because the cross-linking at the N-terminal positions would not impact the open conformation. Therefore, the results suggest that the open-to-closed conformational change induced by cholesterol has a functional implication on SNARE-dependent fusion.

Discussion

Recently, it has been suggested that regulatory lipids such as sterols, diacylglycerol, and phosphoinositides play critical roles in SNARE-dependent fusion as essential cofactors (56, 57). Cholesterol is also shown to be required for some viral fusion systems such as the fusion by α -virus (58). Although the critical importance of the different lipids in various biological fusion events is recognized, the effect of the regulatory lipids on the structures of the fusion proteins is not well understood.

As an initial attempt to uncover the impact of cholesterol on the structures of SNAREs, we investigated the conformational change of the VAMP2 TMD using SDSL EPR. The EPR analysis showed that the TMDs of v-SNARE VAMP2 formed an open scissors-shaped dimer with the pivot point at the N-terminal region in the absence of cholesterol. The overall shape of the dimeric TMD is

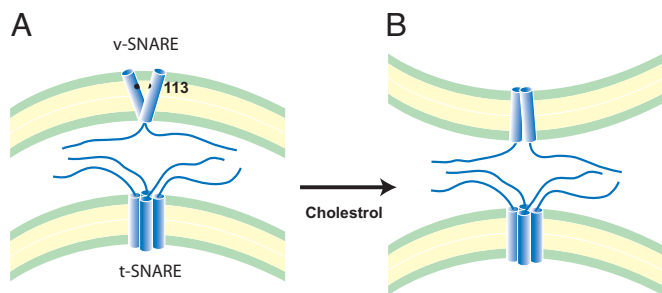


Fig. 5. Hypothetical models for the VAMP2 TMDs in the membrane in the absence (A) and in the presence (B) of cholesterol. In the absence of cholesterol, the dimer has an open-scissors-like conformation that may force the bilayer to have the negative curvature, which is not favorable for fusion. But in the presence of cholesterol, the dimer changes to the parallel conformation, which may not harbor the unfavorable negative curvature. The Syntaxin TMD was modeled as a tetramer based on our earlier work on yeast SNAREs (24).

likely to resemble an inverse wedge that would fit better to the negatively curved bilayer (Fig. 5A). For fusion to occur, however, the bilayers are required to bend in such a way to create positive curvature toward each other. Therefore, the inverse wedge-shaped TMD dimer would act as an inhibitor for such initial bending. Our results show that physiological cholesterol induces a conformational change from the open to the closed, driving the molecular shape to be more favorable for lipid mixing (Fig. 5B). Cholesterol is shown to have the capacity of thickening the bilayer, perhaps because of the rigid nature of its structure (59). We speculate that the closed conformation in the presence of cholesterol reflects the adjustment of the TMD dimer to the thicker bilayer to maintain the hydrophobic match between the bilayer and the TMD (60).

Cholesterol is generally believed to act as a fusion stimulator (61, 62). Its spontaneous negative curvature has the capacity to stabilize the hemifusion state in which the negatively curved surface is dominant. Moreover, it may drive the expansion of the pore, presumably by destabilizing the early fusion pore that is full of an acutely positively curved surface (17). Interestingly, incorporation of 40 mol % cholesterol on the t-SNARE vesicle has less of an effect on the rate of SNARE-mediated lipid mixing than it does for the v-SNARE vesicles. The results raise the possibility that cholesterol impacts membrane fusion not only through its preference for specific membrane curvature but also through its influence on the TMD conformation in the membrane. Our results suggest that the latter is significantly more important for SNARE-mediated membrane fusion and it is predominantly through its influence on the structure of the v-SNARE TMD.

Although it is thought that the TMD of VAMP2 plays a critical role in membrane fusion, controversy exists concerning its structure in the membrane. Computational and mutagenesis studies predicted that the v-SNARE TMD exists as a dimer with 2 helices oriented with a cross-angle of 38° (38). Furthermore, alanine-scanning mutagenesis indicated that residues 99, 102, 103, 107, 110, and 111 are involved in homodimerization of the TMD. In contrast, the infrared (IR) study by Bowen et al. (41) showed that the VAMP2 TMD exists as monomers in the DMPC bilayer with an unusual 35° tilt with respect to the bilayer normal. The EPR-based structural model in the absence of cholesterol might be similar to the structural model proposed by Bowen et al., except that there is a weak and dynamic helix–helix interaction at the N-terminal region. This weak interaction was not detectable even in the sensitive EPR-line shape analysis. Likewise we suspect that this interaction was not detectable in the IR study. Yet it appears that the EPR-based model in the presence of the cholesterol was quite consistent with the model predicted from the computer modeling and the alanine-scanning experiments. In our model, residues 99, 103, 110, and 113 reside at the dimer interface, which were similar to those positions responsive to alanine mutations.

Perhaps, the monomer–dimer controversy stems from the labile equilibrium between monomers and dimers that may be easily shifted by the changes in conditions such as the lipid composition. Provided the uncertainty of the new DEER method to estimate the stoichiometry of the TMD it is still possible that the monomer–dimer equilibrium exists. Therefore, an alternative scenario might be that cholesterol shifts equilibrium toward the dimer, and the dimerization might be the main event that is responsible for the stimulation of SNARE-dependent lipid mixing by cholesterol.

Although the EPR data are consistent with the dimer model, the periodic distance variation along the sequence (Fig. 3C) is much shallower than normally expected for the well-defined dimer. The results indicate that the dimer may not have a well-defined point of symmetry, and the interaction between 2 monomers may be promiscuous despite some preference given to the helical face consisting of residues 99, 103, 106, and 113 (Fig. 3D). The syntaxin TMD is unusual in that it contains as many as 11 β -branched amino acids (Fig. 3D), which may allow neither a well-defined interface between 2 interacting helices nor tight packing, reducing the amplitude of the distance variation significantly along the sequence.

In this work, we used the fluorescence lipid-mixing assays, not the content mixing assay, to investigate the SNARE-dependent membrane fusion. However, the caveats of the lipid-mixing assay are the potential complications arising from liposome instability and leaking (63), lipid flip–flop without fusion (64), and spontaneous lipid dye transfer between apposed membranes (65), warranting further investigations using the content mixing assay which is less vulnerable to these complications.

In summary, we have shown that cholesterol plays a role in modifying the v-SNARE structure in the membrane to be favorable for membrane fusion. Cholesterol induces the conformational change of the v-SNARE TMD dimer, which converts the molecular shape from an open scissors-like to a closed scissors-like shape, thereby modifying the membrane curvature to be favorable for fusion.

Materials and Methods

Plasmid Constructs and Site-Directed Mutagenesis. Full-length VAMP2 (amino acids 1–116) was inserted into the pGEX-KG vector to make the glutathione S-transferase fusion protein. To introduce unique cysteine in the VAMP2 site-specifically, we changed the native cysteine C103 to alanine. QuikChange site-directed mutagenesis was used to make all cysteine mutants. Recombinant Syntaxin 1A without the Habc domain (SynHT, amino acids 168–288) and recombinant SNAP-25, in which 4 palmitoylation sites (cysteines) on the interhelical loop were changed to alanines (amino acids 1–206), were also made as GST fusion proteins. The DNA sequences were confirmed by the Iowa State University sequencing facility.

Protein Expression, Purification, and Spin Labeling. The experiments were carried out as described (30, 35). The labeling efficiency was calculated by dividing the spin concentration with the protein concentration. For all samples, the labeling efficiency was >85%.

Vesicle Preparation and Membrane Reconstitution. Large unilamellar vesicles of a 100-nm diameter were prepared as described in refs. 30 and 35. The lipid-to-protein molar ratio was 200:1 (except as noted). The reconstitution efficiency was estimated by determining the protein concentration before and after reconstitution by using EPR. For all samples, the efficiency was \approx 85%. For the lipid-mixing assay, the mixture of POPC, DOPS, and cholesterol in a molar ratio of 85:15:0, 65:15:20 or 45:15:40 was used. Additionally, NBD-PS and rhodamine-PE (1.5 mol % each) were added to the v-SNARE vesicles for fluorescence detection of lipid mixing. The t-SNARE proteins SynHT and SNAP-25, in a molar ratio of 1:1, were premixed, and the mixture was left at room temperature for \approx 60 min to form the complex before the reconstitution. In all cases the proteins were reconstituted by using the dialysis method described in ref. 66.

Lipid-Mixing Assay. The v-SNARE liposomes were mixed with the t-SNARE liposomes at a molar ratio of 1:9. The total volume of the reaction solution was 100 μ L containing 1 mM lipids. Fluorescence was measured at excitation and emission wavelengths of 465 and 530 nm, respectively. The intensity changes were recorded by the Varian Cary Eclipse model fluorescence spectrophotometer using

a quartz cell of 100 μL with 2-mm path length. All of the lipid-mixing experiments were carried out at 37 $^{\circ}\text{C}$. Approximately 50 min after the reaction, we added 1 μL of 10%-reduced Triton to complete the lipid mixing assay and to obtain the maximum fluorescence intensity (MFI). The inner leaflet-mixing assay was performed following the procedure described in ref. 30.

CW-EPR Data Collection and Analysis. CW-EPR spectra were collected by using the Bruker ESP 300 spectrometer equipped with a loop-gap resonator (Medical Advances) and a low-noise microwave amplifier (Miteq). Room-temperature (293 K) and low-temperature (130 K) spectra were collected for all reconstituted proteins. To measure interspin distances from CW-EPR line shapes, Fourier deconvolution analysis was used (37).

Distance Measurements with the 4-Pulse Ku-band DEER. Four-pulse DEER (44–46) experiments were conducted by using a home-built Ku-band (17.35 GHz) pulse FT-ESR/DQC spectrometer as described (52–54). All measurements were conducted at 65 K by using a 15- μL sample volume. All 3 detection mode pulses

had 20-ns lengths. The pump π -pulse was typically 12–20 ns. The dipolar signals measured by DEER were analyzed by the L-curve Tikhonov regularization method (67) and refined by MEM (68) to recover distance distributions.

Disulfide Cross-Linking and Lipid-Mixing Assay. Wild-type VAMP2 has a single native cysteine at position 103. C103 or mutant Y113C was cross-linked with CuSO_4 (36 μM) and phenanthroline (72 μM) at room temperature for 40 min. Cross-linked dimers C103C-C and Y113C-C and non-cross-linked C103 and Y113C were separately reconstituted into the fluorescent vesicles, whereas the t-SNARE binary complex was inserted into the nonfluorescent vesicle (POPC/DOPS = 85:15 in moles). The lipid-mixing assay was performed by following the procedure described above except that the 1:1 ratio of v-liposome and t-liposome was used this time. The total volume of the reaction solution was 100 μL containing 0.4 mM lipids.

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