

Calcium Promotes the Formation of Syntaxin 1 Mesoscale Domains through Phosphatidylinositol 4,5-Bisphosphate*

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Phosphatidylinositol 4,5-bisphosphate (PI(4,5)P₂) is a minor component of total plasma membrane lipids, but it has a substantial role in the regulation of many cellular functions, including exo- and endocytosis. Recently, it was shown that PI(4,5)P₂ and syntaxin 1, a SNARE protein that catalyzes regulated exocytosis, form domains in the plasma membrane that constitute recognition sites for vesicle docking. Also, calcium was shown to promote syntaxin 1 clustering in the plasma membrane, but the molecular mechanism was unknown. Here, using a combination of superresolution stimulated emission depletion microscopy, FRET, and atomic force microscopy, we show that Ca²⁺ acts as a charge bridge that specifically and reversibly connects multiple syntaxin 1/PI(4,5)P₂ complexes into larger mesoscale domains. This transient reorganization of the plasma membrane by physiological Ca²⁺ concentrations is likely to be important for Ca²⁺-regulated secretion.

Neurotransmitter release requires tight spatial and temporal control. Temporal control is achieved by the interplay between Ca²⁺ influx and synaptotagmin 1, the main Ca²⁺ sensor on the synaptic vesicle (1, 2). Spatial control is achieved by the specific lateral organization of the presynaptic SNARE proteins syntaxin 1 and SNAP25 in the plasma membrane (3–5). Syntaxin 1 and SNAP25 complex with synaptobrevin 2 (VAMP2) in the synaptic vesicle, resulting in membrane fusion and the release

of neurotransmitters (5, 6). It is well established that syntaxin 1 and SNAP25 are not randomly distributed over the plasma membrane of neurons and neuroendocrine cells but form clusters of ~40–100 nm in diameter (7–9). These clusters are necessary for the recruitment of the neurotransmitter-containing vesicles to the plasma membrane (10–12). Clustering of SNAREs has been intensively studied, and different mechanisms affect their lateral organization, including both protein-protein and protein-lipid interactions (3, 4, 8, 12–14). Polyphosphoinositides belong to the components shown to be important for syntaxin domain formation. Syntaxin 1 contains a polybasic stretch juxtaposed to its transmembrane domain, which interacts electrostatically with polyphosphoinositides (15–17), including PI(4,5)P₂,³ which represents more than 80% of total lipids in syntaxin 1 domains (16).

Apart from triggering synaptic vesicle release, calcium ions induce a reorganization of the plasma membrane, resulting in larger, mesoscale domains of SNAREs, including syntaxin 1 (18). This clustering relates to the net charge of the protein, with more anionic proteins forming more pronounced clusters with Ca²⁺, arguing for a charge bridging effect. In addition, it is well established that Ca²⁺, but not Mg²⁺, also induces domain formation of PI(4,5)P₂ by means of charge bridging, which results in connection between multiple PI(4,5)P₂ molecules (19–23). Because the polybasic juxtamembrane stretch of syntaxin 1 interacts with the multiple negative charges in the headgroup of PI(4,5)P₂ (15–17), we hypothesized that Ca²⁺ may cluster syntaxin 1 indirectly via PI(4,5)P₂. This would result in coalescence of multiple smaller syntaxin 1/PI(4,5)P₂ clusters into a larger domain. Indeed, using a combination of superresolution stimulated emission depletion (STED) nanoscopy, FRET, and atomic force microscopy (AFM), we now show that, under physiological conditions, Ca²⁺ ions can act as a charge bridge and induce the formation of syntaxin 1/PI(4,5)P₂ domains at the mesoscale.

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³ The abbreviations used are: PI(4,5)P₂, phosphatidylinositol 4,5-bisphosphate; STED, stimulated emission depletion; AFM, atomic force microscopy; Fmoc, N-(9-fluorenyl)methoxycarbonyl; DOPC, 1,2-dioleoyl-sn-glycero-3-phosphatidylcholine; LUV, large unilamellar vesicle; FWHM, full width at half-maximum intensity; TMD, transmembrane domain.

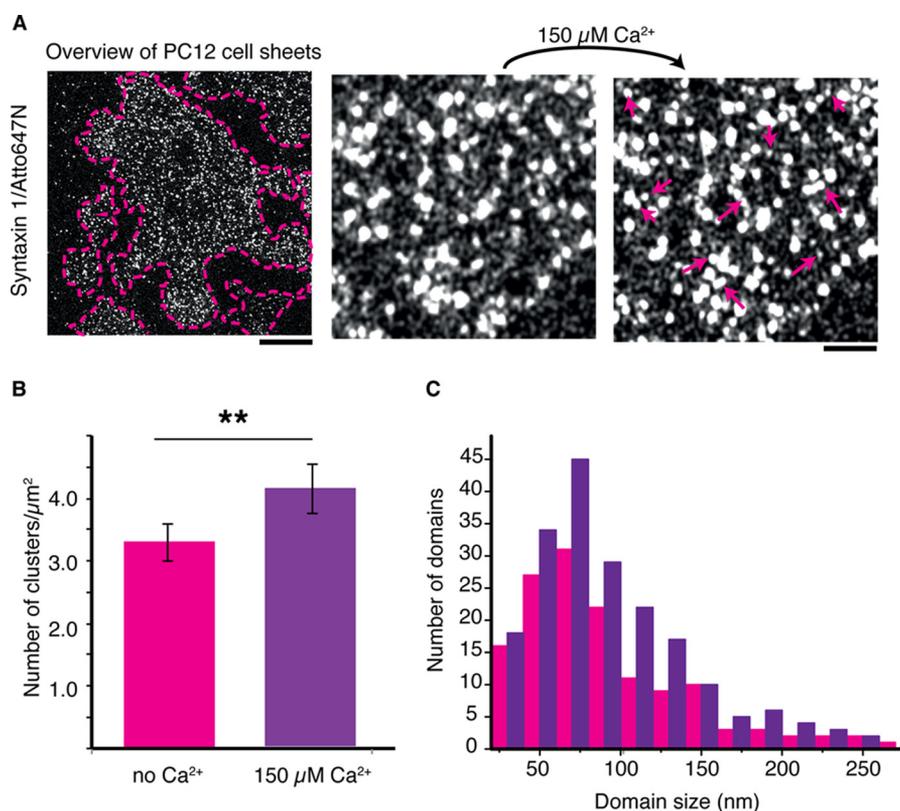


FIGURE 1. Calcium promotes clustering of syntaxin 1 in the plasma membrane of PC12 cells. *A*, overview of a typical PC12 membrane sheet immunostained for syntaxin 1 and imaged by STED (*left panel*, scale bar = 4 μm) and sections of the sheet at larger magnification both before (*center panel*) and after (*right panel*, scale bar = 1 μm) addition of 150 μM Ca^{2+} . Newly appeared syntaxin domains after addition of Ca^{2+} are indicated by arrows. *B*, the density of syntaxin 1 clusters increased by $\sim 25\%$ after addition of Ca^{2+} . Error bars indicate the range from three independent experiments with at least 10 sheets analyzed. **, $p < 0.01$, two-sided paired t test. *C*, the size distributions (FWHM) of syntaxin 1 domains in the plasma membrane in the absence (*pink*) and presence (*purple*) of 150 μM Ca^{2+} . Data are pooled from three independent experiments.

Experimental Procedures

Materials—Syntaxin 1 TMD (residues 266–288; sx-1 TMD *Rattus norvegicus* sequence) and syntaxin 1 TMD mutant (sx-1 TMD with the following mutations: K265A and K266A) were synthesized using Fmoc solid-phase synthesis as described in Ref. 16. The fluorescent dyes Atto647N NHS-ester (Atto-Tec) and Rhodamine red succinimidyl ester (Life Technologies) were coupled to the N termini of the peptides during the Fmoc synthesis.

DOPC (1,2-dioleoyl-*sn*-glycero-3-phosphatidylcholine), 1,2-dioleoyl-*sn*-glycero-3-phosphatidylserine, and PI(4,5)P₂ (1,2-dioleoyl-*sn*-glycero-3-phosphatidyl-(1'-myo-inositol-4',5'-bispophosphate)) were purchased from Avanti Polar Lipids. Atto647N labeled at the SN1 position of PI(4,5)P₂ and Atto590 coupled to ceramide were gifts from Dr. Vladimir Belov (MPI-BPC, Göttingen, Germany).

Cell Analyses—We used the pheochromocytoma cell line PC12 from *R. norvegicus* (24) to prepare native membrane sheets by gentle sonication as described in Ref. 12. Sonication buffer contained 20 mM potassium HEPES (pH 7.4), 120 mM potassium gluconate, 20 mM potassium acetate, 2 mM ATP, and 0.5 mM DTT. The antibody used for immunohistochemistry was anti-syntaxin 1 mouse IgG1 (Sigma, clone HPC-1) labeled with KK114-maleimide (a gift from Dr. Vladimir Belov, Max Planck Institute for Biophysical Chemistry).

FRET Measurements—For FRET measurements, we prepared large unilamellar vesicles (LUVs) that contained Rhoda-

mine Red coupled to sx-1 TMD (donor fluorophore) and Atto647N coupled to sx-1 TMD (acceptor) as described in Ref. 25. The total protein-to-lipid molar ratio in our FRET measurements was 1:1000. Excitation was at 560 nm, and emission was collected from 570–700 nm, with 1-nm slit widths on a FluoroMax-2 fluorescence spectrometer (Horiba). We corrected for cross-talk resulting from acceptor excitation using samples containing only the acceptor fluorophore. The FRET efficiency was calculated as the ratio of emission intensities at 660 nm (acceptor maximum) over 580 nm (donor maximum).

STED Nanoscopy—For STED nanoscopy, a home-built setup was used, with pulsed excitation lasers at 595 and 640 nm. The fluorescence was collected from 600–640 and 660–720 nm with avalanche photo diodes (Excelitas and Micro Photon Devices). Superresolution was achieved using a STED laser (775 nm, 20 MHz pulsed fiber laser, IPG Photonics). By combining a 2π vortex phase plate (RPC Photonics) and a $\lambda/4$ plate, the typical “donut-shaped” focal intensity distribution of the STED beam was produced. Using the same STED beam for both dyes inherently ensured a colocalization accuracy far below the resolution limit (26). Another setup employed was a commercially available two-color STED setup (Abberior Instruments, Göttingen, Germany). This setup had two pulsed excitation lasers at 594 and 640 nm and a pulsed STED laser at 775 nm. The setup had a QUAD beam scanner (Abberior Instruments). Pulse energies ranging from 3–8 nJ in the back aperture of the objective yielded a lateral resolution of down to 30 nm. Data

Calcium Clusters Syntaxin 1 through PI(4,5)P₂

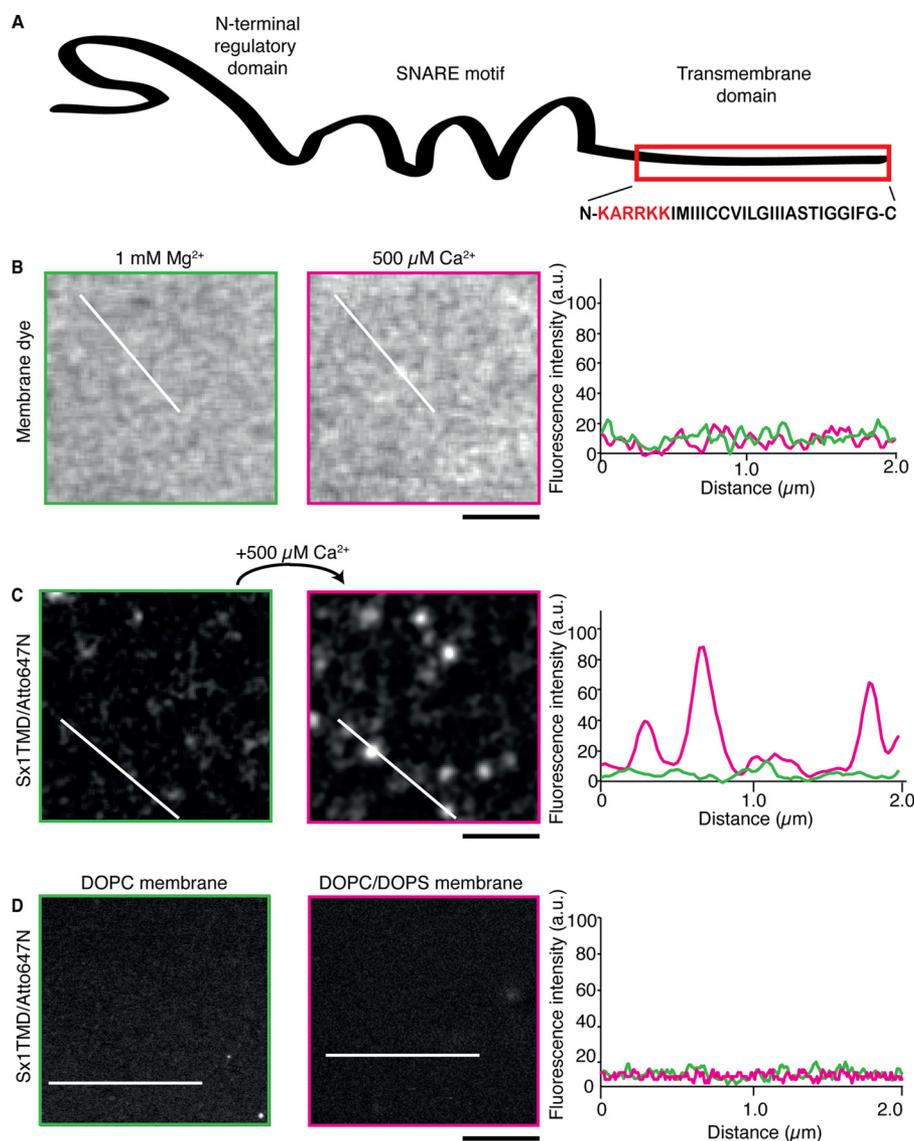


FIGURE 2. Calcium-promoted syntaxin clustering requires PI(4,5)P₂. *A*, domain structure of syntaxin 1. The polybasic stretch juxtaposed to the transmembrane domain is marked in red. *B*, two-color STED images of glass-supported membranes containing sx-1 TMD (1:10,000 protein:lipid ratio) and composed of 97 mol% DOPC and 3 mol% PI(4,5)P₂. Ceramide-Atto594 was used as a lipid marker (membrane dye). Buffers contained 1 mM Mg²⁺ (left panel, green in the intensity profile) or 500 μM Ca²⁺ (center panel, magenta in the intensity profile). Fluorescence intensity profiles along the cross-sections are indicated in the right panel in absolute units. *C*, the same as *B* but with sx-1 TMD labeled with Atto647N. *D*, the same as *B* but with membranes composed of pure DOPC (left panel) or 80 mol% DOPC and 20 mol% 1,2-dioleoyl-*sn*-glycero-3-phosphatidylserine (DOPS, center panel), both in the presence of 500 μM Ca²⁺. Representative data from multiple independent experiments are shown. Fluorescence intensity profiles along the cross-sections are indicated in the right panel in counts. Scale bar = 1 μm.

acquisition was done using ImSpector software. The density of clusters was analyzed using the particle analysis plugin in the Fiji software, and the cluster correlation was obtained using Pearson's correlation analysis from the Fiji software tools (27). Cluster sizes were calculated as the full width at half-maximum intensities (FWHM).

AFM Imaging of Stacked Lipid Bilayers—Glass coverslips were cleaned using a Plasma cleaner Fempto timer with a 40-kHz, 100-W generator (Diener Electronic), and a lipid/sx-1 TMD bilayer was generated by spin-coating as described in Ref. 25. The reconstituted bilayers were imaged with a Cervantes full mode AFM system (Nanotec) using AC40TS cantilevers ($f_0 = 110$ kHz, $k = 0.1$ N/m, Olympus) as described previously (28). Calibration of the cantilevers was accomplished by using the thermal noise spectrum. We employed the Jumping Mode

Plus (jump-off, 100 nm; sample points, 50), which allows scanning at controlled vertical forces between 0.2 nN and several nanonewtons (29).

Results

We first confirmed the previously reported finding (18) that elevated Ca²⁺ promotes clustering of syntaxin 1 in the plasma membrane. We employed PC12 cell sheets, which are a widely used model system for studying the lateral organization of membranes (8, 9, 12, 13, 16, 25). Using STED nanoscopy, we obtained high-resolution images of PC12 plasma membranes immunolabeled for syntaxin 1 before and after the addition of 150 μM Ca²⁺ (Fig. 1A). After analyzing at least 10 cell sheets from three different experiments, we observed that the average cluster density of syntaxin 1 increased from 3.3 ± 0.3 clusters

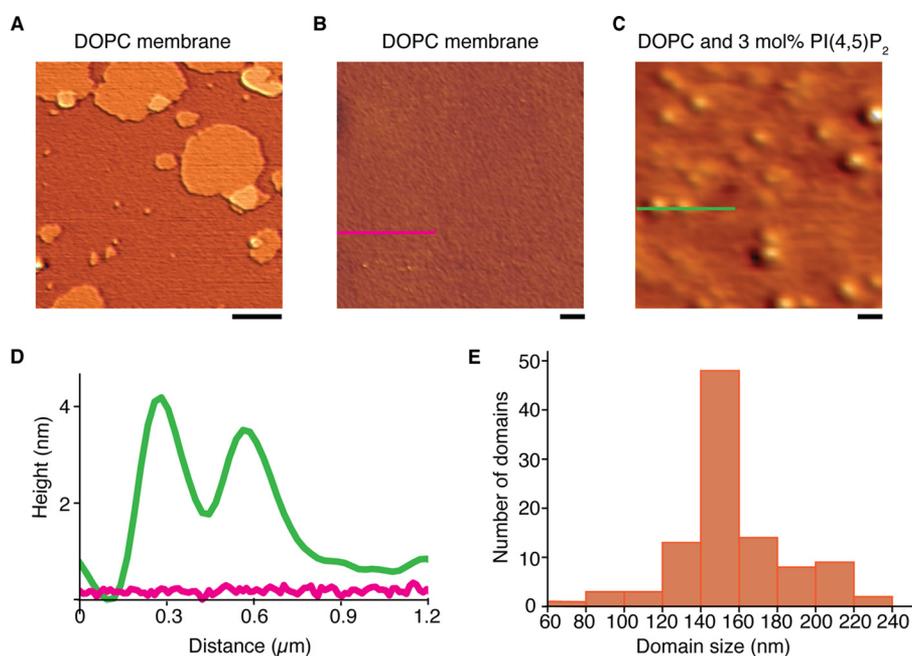


FIGURE 3. **Sx-1 TMD/PI(4,5)P₂ domains reconstituted in lipid bilayers observed by AFM.** A, representative AFM image of stacked lipid bilayers on a glass support. Lipid bilayers were composed of pure DOPC and sx-1 TMD (1:10,000 protein:lipid ratio) in the presence of 500 μM Ca^{2+} . Note that the membrane had patches of multiple layers on top of each other. Scale bar = 2 μm . B, magnification of the membrane region in A. Scale bar = 0.3 μm . C, the same as B but with bilayers containing 3 mol% PI(4,5)P₂. D, height profiles along the cross-sections indicated in B (pink) and C (green). E, size distribution of sx-1 TMD domains from PI(4,5)P₂-containing membranes (FWHM). Data are pooled from at least 10 independent reconstitutions.

μm^{-2} to 4.1 ± 0.4 clusters μm^{-2} upon calcium addition (Fig. 1B). In addition to this increase in density, Ca^{2+} increased the size of the domains from an average diameter of ~ 90 nm in the absence of Ca^{2+} to ~ 105 nm after Ca^{2+} addition (FWHM, Fig. 1C). Assuming circular domains, this equates to an $\sim 40\%$ increase in domain area upon calcium addition, although this increase is an underestimate because domain sizes are overestimated by ~ 5 – 10 nm because of the antibody staining. At our STED resolution, the sizes of the antibodies will contribute substantially to the measured domain sizes (the so-called “umbrella effect”) (7).

We then delineated the role of calcium on syntaxin 1 clustering in precisely controllable model membranes. We used fluorescently labeled syntaxin 1 TMD peptide (sx-1 TMD, residues 257–288). Apart from the polybasic linker region, this peptide does not contain the cytosolic domains (Fig. 2A), which allowed for exclusion of any contribution of cytosolic protein-protein interactions on syntaxin clustering. Our lipid mixtures for membrane reconstitutions contained DOPC without cholesterol. We recently showed that DOPC bilayers have a thickness of ~ 3.5 nm, which matches the hydrophobic length of sx-1 TMD, and that this membrane system displays minimal clustering caused by hydrophobic mismatching (25). This model system therefore allowed us to focus on the effect of calcium on interactions between sx-1 TMD and PI(4,5)P₂. In stacked lipid bilayers, the addition of Ca^{2+} did not affect the lipid bilayer structure as visualized by the fluorescently labeled lipid analogue ceramide-Atto590 (Fig. 2B). However, 500 μM of Ca^{2+} caused the clustering of reconstituted sx-1 TMD, provided PI(4,5)P₂ was present in the membrane (Fig. 2C). We did not observe clustering of syntaxin in membranes composed of only DOPC or a mixture of 80 mol% DOPC and 20 mol% 1,2-dioleoyl-*sn*-glycero-3-phosphatidylserine (Fig. 2D).

To test whether the observed clusters were representing lateral membrane domains or small membrane vesicles on top of the bilayer, we used AFM. We prepared supported lipid bilayers with sx-1 TMD on plasma-treated glass surfaces (Fig. 3A). When the membrane was composed of only DOPC, scanning showed a homogeneously flat surface regardless of the presence of calcium (Fig. 3, B and D). In contrast, and comparable with our fluorescence data, clusters were clearly observed when the membranes contained 3 mol% PI(4,5)P₂, provided calcium was present (Fig. 3, C and D). Here the circular sx-1 TMD domains had an average size of 157 ± 2.7 nm (full width at half-maximum height, Fig. 3E), which is similar to the size determined by STED microscopy. In the absence of calcium, we did not observe clustering of syntaxin 1, regardless of the presence or absence of PI(4,5)P₂. The average height of the domains was only 4 nm. Although this height might be taller than the expected dimensions of sx-1 TMD alone (~ 2 nm), possibly because of buckling of the membrane, this height is far too small for any membrane vesicle. These experiments demonstrate that calcium can cluster sx-1 TMD in the presence of PI(4,5)P₂ and supports our hypothesis.

Next we tested the specificity and reversibility of calcium-triggered sx-1 TMD domain formation. To this end, we reconstituted sx-1 TMD in membranes that contained 3 mol% PI(4,5)P₂ and recorded a series of STED images of the same membrane regions while changing the components of the buffer (Fig. 4). The addition of Mg^{2+} at a final concentration of 1 mM did not cause any clustering of the sx-1 TMD. However, the addition of 500 μM Ca^{2+} immediately triggered the formation of sx-1 TMD clusters with sizes between 70 and 200 nm (Fig. 4B). These domains were dependent on the presence of Ca^{2+} ions because chelating calcium with 0.5 M EGTA fully reversed sx-1 TMD clustering. The addition of 500 μM Ca^{2+} also triggered clustering of a

Calcium Clusters Syntaxin 1 through PI(4,5)P₂

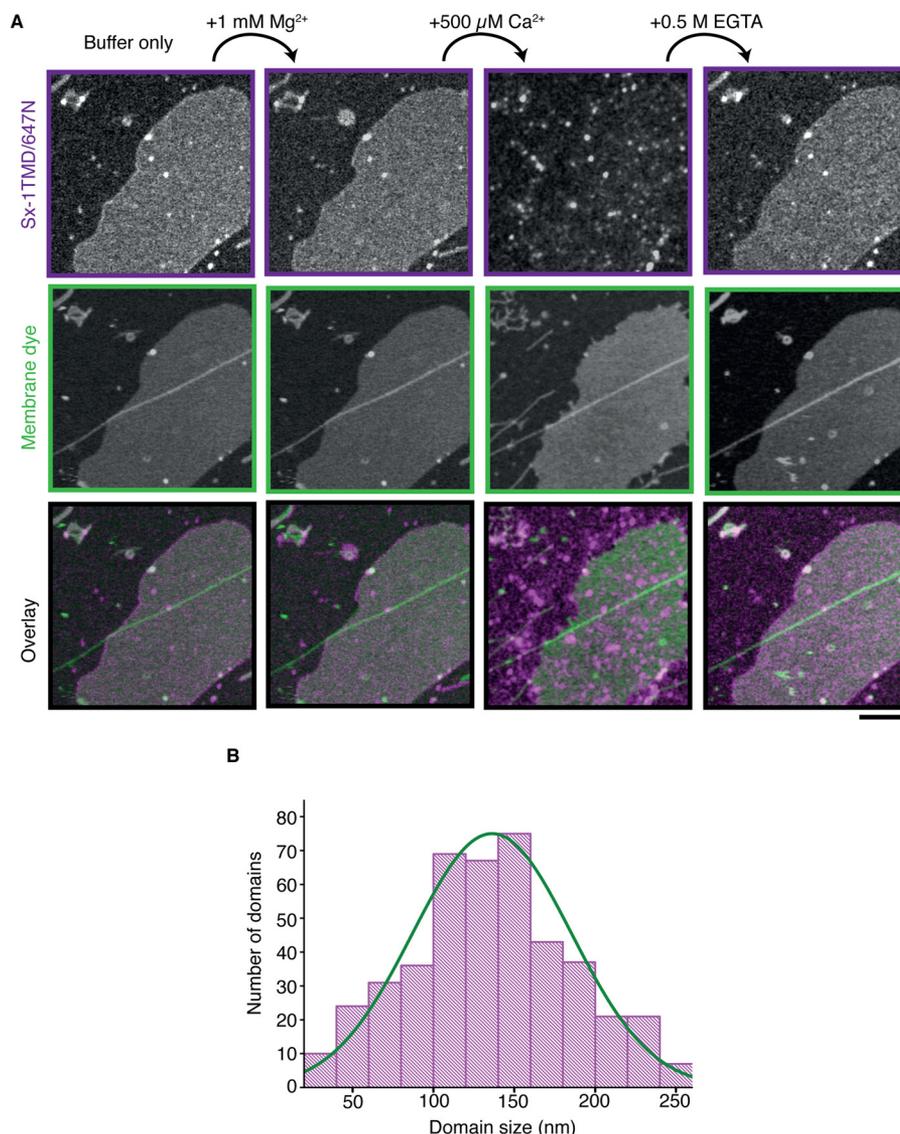


FIGURE 4. Reversible clustering of syntaxin 1/PI(4,5)P₂ domains induced by Ca²⁺. *A*, representative series of STED images of membranes composed of 97 mol% DOPC and 3 mol% PI(4,5)P₂ with sx-1 TMD reconstituted (protein:lipid ratio of 1:10,000, *magenta* in the overlay) and ceramide labeled with Atto590 (*green*, membrane dye). Mg²⁺ (1 mM), Ca²⁺ (500 μM), and EGTA (0.5 M) were added sequentially, and the same membrane area was imaged after each addition. Scale bar = 4 μm. *B*, size distribution of sx-1 TMD domains in the presence of 500 μM Ca²⁺ (FWHM, data from three independent reconstitutions).

PI(4,5)P₂ variant containing an acyl chain labeled with Atto647N (Fig. 5), demonstrating that Ca²⁺ induced clustering not only of sx-1 TMD but also of PI(4,5)P₂.

To further characterize the molecular interactions between sx-1 TMD and the polar headgroup of PI(4,5)P₂, we employed a recently developed FRET-based assay (15, 25). We reconstituted sx-1 TMD in LUVs with half of the peptide labeled with Rhodamine Red (FRET donor fluorophore) and the other half with Atto647N (acceptor fluorophore), both conjugated to the N-terminal end of the peptide. We measured the emission spectra in samples before and after the addition of 150 μM Ca²⁺. As expected, the polybasic juxtamembrane stretch of sx-1 TMD interacted with PI(4,5)P₂, resulting in protein clustering, and this interaction was significantly increased after the addition of Ca²⁺ (Fig. 6A). To confirm the specific interaction of the polybasic juxtamembrane stretch of sx-1 TMD with PI(4,5)P₂, we mutated two lysine residues located in the polybasic stretch to

neutral alanines (K264A and K266A). Mutating these two lysines disrupts the interaction of syntaxin 1 with PI(4,5)P₂ (15, 16). Indeed, this sx-1 TMD mutant showed reduced clustering with PI(4,5)P₂, and the Ca²⁺ effect was also diminished. Sx-1 TMD clustering was also reduced in the absence of polyvalent PI(4,5)P₂, and this clustering could not be rescued by monovalent phosphatidylserine (one negative charge, Fig. 6A).

In the final set of experiments, we investigated whether polyvalent ions other than calcium can decrease the electrostatic interactions responsible for syntaxin clustering by charge screening. To this end, we included both Mg²⁺ and ATP, which are present in the cytoplasm, at relatively high concentrations of 0.5–5 mM and 1–2 mM, respectively (30, 31). Indeed, in the absence of Ca²⁺, decreased clustering of sx-1 TMD was observed when 5 mM Mg²⁺ and/or 5 mM ATP was included in the buffer (Fig. 6B). However, Ca²⁺ was able to overcome this charge screening effect and increased sx-1 TMD clustering.

Taken together, we conclude that Ca²⁺ can act as a charge bridge that merges multiple small sx-1 TMD/PI(4,5)P₂ clusters into larger membrane domains.

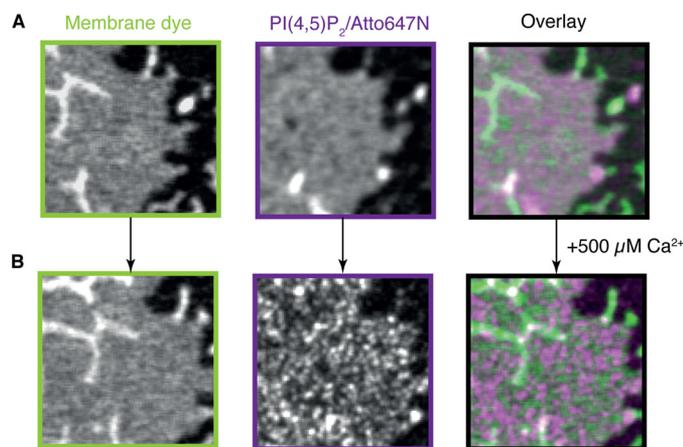


FIGURE 5. Ca²⁺ induces PI(4,5)P₂ clustering independently of sx-1 TMD. STED images of membrane composed of 97 mol% DOPC, 3 mol% unlabeled PI(4,5)P₂, 0.1 mol% PI(4,5)P₂ labeled with Atto647N (magenta in the overlay), and 0.1 mol% DOPE-OG (Oregon green labeled, 1,2-dioleoyl-*sn*-glycero-3-phosphatidylethanolamine; green, membrane dye). *A* and *B*, the same membrane area incubated in the absence and presence of 500 μM Ca²⁺, respectively. Representative images from multiple independent experiments are shown. Scale bar = 1 μm.

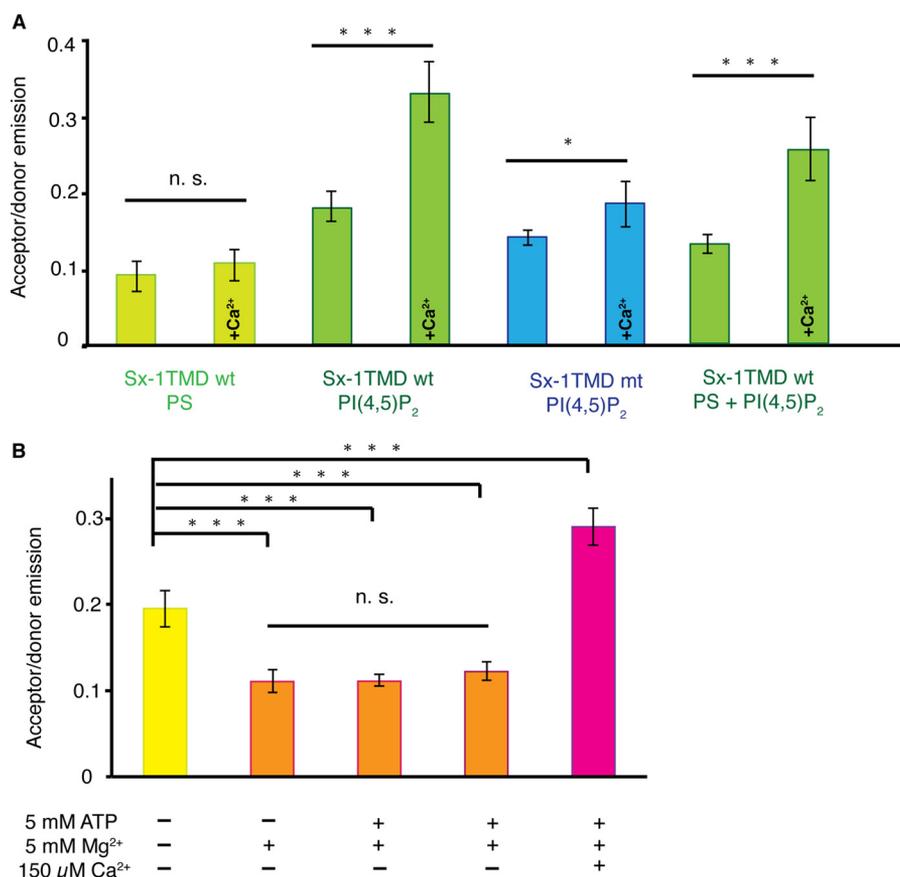


FIGURE 6. Calcium induces coalescence of multiple syntaxin 1/PI(4,5)₂ complexes. *A*, membrane clustering by FRET on LUVs containing sx-1 TMD labeled with Rhodamine Red (donor fluorophore) and Atto647N (acceptor fluorophore). 150 μM Ca²⁺ significantly increased the FRET efficiency in the presence of 3 mol% PI(4,5)P₂ but not in LUVs that contained 20 mol% 1,2-dioleoyl-*sn*-glycero-3-phosphatidylserine in the absence of PI(4,5)P₂. Mutation of the sx-1 TMD (K264A, K265A; *Sx-1 TMD mt*) significantly reduced oligomerization both in the presence and absence of Ca²⁺ compared with the wild-type (wt). *B*, charge screening by ATP and Mg²⁺. LUVs were prepared as in *A*, and FRET was measured in buffer that was supplemented with 5 mM Mg²⁺, 5 mM ATP (disodium salt), or both and in the absence or presence of 150 μM Ca²⁺. Error bars indicate the range from three independent reconstitutions. ***, *p* < 0.001; *, *p* < 0.05; n.s., not significant; two-sided, paired *t* test).

Calcium Clusters Syntaxin 1 through PI(4,5)P₂

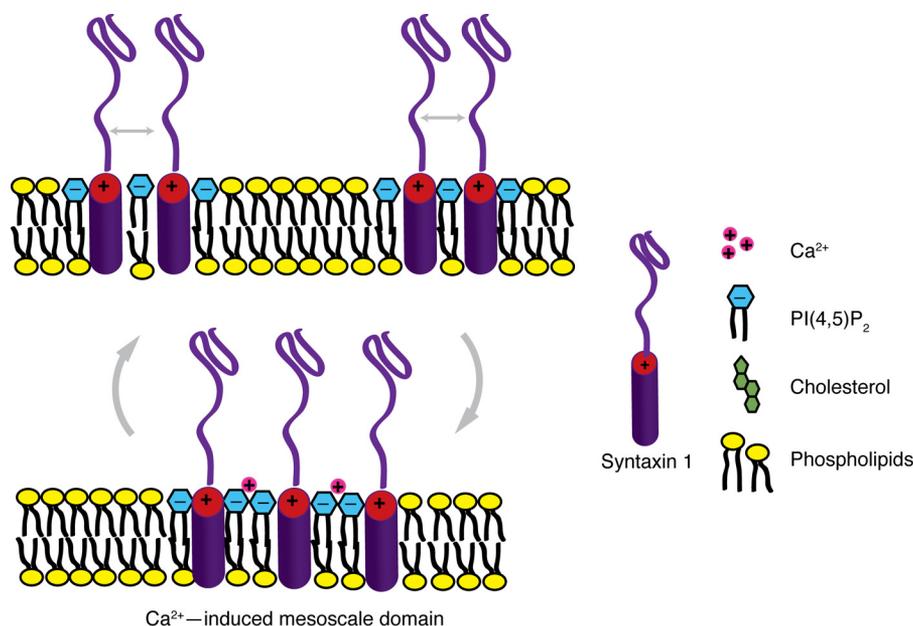


FIGURE 7. **Model of calcium-induced syntaxin domain formation.** Ca²⁺ specifically bridges PI(4,5)P₂ molecules and induces the coalescence of syntaxin-1/PI(4,5)P₂ clusters into larger mesoscale domains.

fruit fly, syntaxin clustering at synaptic boutons is governed by phosphatidylinositol 3,4,5-triphosphate (17). Accordingly, not only syntaxin 1, but also other SNAREs, can interact with PI(4,5)P₂. For instance, syntaxin 4 and the R-SNARE synaptobrevin 2 contain polybasic stretches adjacent to their TMDs that were shown to bind to PI(4,5)P₂ (25, 33). Removal of these positive residues in synaptobrevin 2 inhibits exocytosis (33). It remains to be established whether synaptobrevin 2 can also be driven into clusters and, if so, whether such clustering is confined to the synaptobrevin pool in the plasma membrane or whether it also occurs in the membrane of secretory vesicles that are devoid of PI(4,5)P₂.

Third, the electrostatic interaction between the polybasic juxtamembrane stretch of syntaxin and the polyanionic headgroup of PI(4,5)P₂ are reduced by other polyvalent ions such as Mg²⁺ and ATP. The fact that Ca²⁺, at a much lower concentration, can overcome this electrostatic shielding shows that clustering is driven not only by bulk electrostatics but specifically involves a defined recognition of the participating molecules (Fig. 7). In PC12 cells, Ca²⁺ promotes clustering not only of syntaxin 1 but also of other SNAREs and integral and peripheral membrane proteins (18). Because many proteins directly or indirectly interact with PI(4,5)P₂, including proteins that organize the cortical cytoskeleton (reviewed in Ref. 32, 34), Ca²⁺-induced PI(4,5)P₂ clustering may well explain these structural rearrangements.

In this study, we observed increased clustering of syntaxin 1 at calcium concentrations of ~500 μM in the microscopy-based experiments. In the more sensitive FRET assay using LUVs, clustering was already observed at lower calcium concentrations of ~150 μM. These concentrations are higher than the global calcium concentration in synaptic boutons, which increases from less than 0.2 μM under resting conditions to ~30 μM after membrane depolarization (35). However, much higher local calcium concentrations (several hundred micromolar) can be reached in the vicinity of synaptic calcium channels upon depolar-

ization (36, 37). Syntaxin 1 domains are located in close proximity to calcium channels (38, 39), and these channels even physically interact with SNAREs (36), suggesting that Ca²⁺-dependent clustering does occur under physiological conditions.

What may be the functional significance of Ca²⁺-induced clustering of syntaxin? Ca²⁺/PI(4,5)P₂-dependent clustering of syntaxin may play a role in exocytosis. We have recently shown that syntaxin 1/PI(4,5)P₂ domains represent the preferred binding sites for the exocytotic calcium sensor synaptotagmin 1 (40, 41). This raises the possibility that a Ca²⁺-induced local increase in syntaxin density promotes SNARE complex formation at the site of release. Alternatively, syntaxin clustering by Ca²⁺ may remove excess SNAREs from the fusion site, thereby preventing hindrance of exocytosis by molecular crowding (5). A third possibility may be that clustering facilitates endocytosis. Ca²⁺-induced mesodomains may help to segregate plasma membrane proteins from the membrane proteins that are destined for endocytosis (42), thereby contributing to the rapid and high-fidelity recycling of synaptic vesicles. The main conclusion from this study is that ionic surface interactions between cations and polyanionic membrane lipids refine the lateral organization of the plasma membrane proteins, and this likely has implications for intracellular membrane trafficking.

Author Contributions—D. M., A. H., G. v. d. B., and R. J. conceived and designed the experiments. M. P. and I. A. T. S. performed the AFM measurements. M. J. and U. D. synthesized the peptides. D. M. performed all other experiments and analyzed the data. D. M., G. v. d. B., and R. J. wrote the paper. All authors reviewed the results and approved the final version of the manuscript.

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Calcium Clusters Syntaxin 1 through PI(4,5)P₂

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