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

Phosphatidylinositol 4,5-bisphosphate (PI(4,5)P$_2$) 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)P$_2$ 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 Ca$^{2+}$ acts as a charge bridge that specifically and reversibly connects multiple syntaxin 1/PI(4,5)P$_2$ complexes into larger mesoscale domains. This transient reorganization of the plasma membrane by physiological Ca$^{2+}$ concentrations is likely to be important for Ca$^{2+}$-regulated secretion.

Neurotransmitter release requires tight spatial and temporal control. Temporal control is achieved by the interplay between Ca$^{2+}$ influx and synaptotagmin 1, the main Ca$^{2+}$ 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). Polysphosphoinositides belong to the components shown to be important for syntaxin domain formation. Syntaxin 1 contains a polyanionic stretch juxtaposed to its transmembrane domain, which interacts electrostatically with polysphosphoinositides (15–17), including PI(4,5)P$_3$, 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 Ca$^{2+}$, arguing for a charge bridging effect. In addition, it is well established that Ca$^{2+}$, but not Mg$^{2+}$, also induces domain formation of PI(4,5)P$_2$ by means of charge bridging, which results in connection between multiple PI(4,5)P$_2$ molecules (19–23). Because the polyanionic juxtamembrane stretch of syntaxin 1 interacts with the multiple negative charges in the headgroup of PI(4,5)P$_2$ (15–17), we hypothesized that Ca$^{2+}$ may cluster syntaxin 1 indirectly via PI(4,5)P$_2$. This would result in coalescence of multiple smaller syntaxin 1/PI(4,5)P$_2$ 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, Ca$^{2+}$ ions can act as a charge bridge and induce the formation of syntaxin 1/PI(4,5)P$_2$ domains at the mesoscale.

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1 To whom correspondence may be addressed: Dept. of Neurobiology, Max Planck Institute for Biophysical Chemistry, 37077 Gottingen, Germany, the
2 To whom correspondence may be addressed: Dept. of Tumor Immunology, Radboud University Medical Center, Geert Grooteplein 10, 6525 GA Nijmegen, The Netherlands. E-mail: rjahn@gwdg.de.

The abbreviations used are: PI(4,5)P$_2$, phosphatidylinositol 4,5-bisphosphate; STED, stimulated emission depletion; AFM, atomic force microscopy; Fmoc, N-(9-fluorenylmethoxycarbonyl; DOPC, 1,2-dioleoyl-sn-glycero-3-phosphatidylcholine; LUV, large unilamellar vesicle; FWHM, full width at half-maximum intensity; TMD, transmembrane domain.
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-(1′/H11032-myoinositol-4′/H11032,5′/H11032-bisphosphate)) 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) and Rhodamine red succinimidyl ester (Life Technologies) were coupled to the N termini of the peptides during the Fmoc synthesis.

FRET Measurements—For FRET measurements, we prepared large unilamellar vesicles (LUVs) that contained Rhodamine 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 \( 2\pi \) 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

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 \( \mu \)m) and sections of the sheet at larger magnification both before (center panel) and after (right panel, scale bar = 1 \( \mu \)m) addition of 150 \( \mu \)M Ca\(^{2+}\). Newly appeared syntaxin domains after addition of Ca\(^{2+}\) are indicated by arrows. B, the density of syntaxin 1 clusters increased by \(~25\%\) 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 \( \mu \)M Ca\(^{2+}\). Data are pooled from three independent experiments.

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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 Femto 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 Ca$^{2+}$ 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 Ca$^{2+}$ (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.
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In stacked lipid bilayers containing 3 mol% PI(4,5)P₂, the addition of 500 mM Ca²⁺ upon calcium addition (Fig. 3B). In addition to this increase in density, Ca²⁺ increased the size of the domains from an average diameter of 90 nm in the absence of Ca²⁺ to 105 nm after Ca²⁺ addition (FWHM, Fig. 1C). Assuming circular domains, this equates to an ∼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 Ca²⁺ did not affect the lipid bilayer structure as visualized by the fluorescently labeled lipid analogue ceramide-Atto590 (Fig. 2B). However, 500 μM of Ca²⁺ 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-glycerol-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 homogenously 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 ± 27 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 Mg²⁺ at a final concentration of 1 mM did not cause any clustering of the sx-1 TMD. However, the addition of 500 μM Ca²⁺ 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 Ca²⁺ ions because chelating calcium with 0.5 M EGTA fully reversed sx-1 TMD clustering. The addition of 500 μM Ca²⁺ also triggered clustering of a
PI(4,5)P$_2$ variant containing an acyl chain labeled with Atto647N (Fig. 5), demonstrating that Ca$^{2+}$ induced clustering not only of sx-1 TMD but also of PI(4,5)P$_2$.

To further characterize the molecular interactions between sx-1 TMD and the polar headgroup of PI(4,5)P$_2$, 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 Ca$^{2+}$. As expected, the polybasic juxtamembrane stretch of sx-1 TMD interacted with PI(4,5)P$_2$, resulting in protein clustering, and this interaction was significantly increased after the addition of Ca$^{2+}$ (Fig. 6A). To confirm the specific interaction of the polybasic juxtamembrane stretch of sx-1 TMD with PI(4,5)P$_2$, 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)P$_2$ (15, 16). Indeed, this sx-1 TMD mutant showed reduced clustering with PI(4,5)P$_2$, and the Ca$^{2+}$ effect was also diminished. Sx-1 TMD clustering was also reduced in the absence of polyvalent PI(4,5)P$_2$, and this clustering could not be rescued by monovalent phosphatidylinerine (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 Mg$^{2+}$ 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 Ca$^{2+}$, decreased clustering of sx-1 TMD was observed when 5 mM Mg$^{2+}$ and/or 5 mM ATP was included in the buffer (Fig. 6B). However, Ca$^{2+}$ was able to overcome this charge screening effect and increased sx-1 TMD clustering.

FIGURE 4. Reversible clustering of syntaxin 1/PI(4,5)P$_2$ domains induced by Ca$^{2+}$. A, representative series of STED images of membranes composed of 97 mol% DOPC and 3 mol% PI(4,5)P$_2$ with sx-1 TMD reconstituted (protein:lipid ratio of 1:10,000, magenta in the overlay) and ceramide labeled with Atto590 (green, membrane dye). Mg$^{2+}$ (1 mM), Ca$^{2+}$ (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 Ca$^{2+}$ (FWHM, data from three independent reconstitutions).
Taken together, we conclude that Ca\(^{2+}\) can act as a charge bridge that merges multiple small sx-1 TMD/PI(4,5)P\(_2\) clusters into larger membrane domains.

**Discussion**

In this study, we demonstrate that Ca\(^{2+}\) induces the coalescence of syntaxin-1/PI(4,5)P\(_2\) clusters into larger mesoscale domains. Three main conclusions can be drawn from our findings. First, calcium only promotes clustering of the sx-1TMD construct in the presence of PI(4,5)P\(_2\). This corroborates our previous findings showing that PI(4,5)P\(_2\) is essential for clustering of syntaxin 1 and that targeting of the phosphatase domain of synaptojanin 1 (a PI(4,5)P\(_2\) phosphatase) to the plasma membrane causes the dispersion of syntaxin 1 clusters in the plasma membrane of PC12 cells (16).

Second, only Ca\(^{2+}\) but not Mg\(^{2+}\) promotes membrane clustering of syntaxin 1/PI(4,5)P\(_2\). This finding correlates well with several studies showing that Ca\(^{2+}\) specifically induces PI(4,5)P\(_2\) domains (19–23). As explained in these studies, the Ca\(^{2+}\) specificity is due to the charge density distributions and the matching of chelating properties between Ca\(^{2+}\) and the polynegative headgroup of PI(4,5)P\(_2\). Although in mammalian cells PI(4,5)P\(_2\) is the dominant phosphoinositide species in the plasma membrane (32) and is present at high concentrations in syntaxin 1 clusters (16), we do not expect that Ca\(^{2+}\)-promoted clustering of syntaxin 1 is specific for PI(4,5)P\(_2\). Not only can calcium cluster other polyphosphoinositide species as well (19), but in...
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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 micromolars) can be reached in the vicinity of synaptic calcium channels upon depolarization (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.

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References

work is a major factor influencing the organization of the secretory machinery in chromaffin cells. J. Cell Sci. 124, 727–734