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Models of synaptotagmin-1 to trigger Ca^{2+} -dependent vesicle fusion

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Vesicles in neurons and neuroendocrine cells store neurotransmitters and peptide hormones, which are released by vesicle fusion in response to Ca^{2+} -evoking stimuli. Synaptotagmin-1 (Syt1), a Ca^{2+} sensor, mediates ultrafast exocytosis in neurons and neuroendocrine cells. After vesicle docking, Syt1 has two main groups of binding partners: anionic phospholipids and the SNARE (soluble *N*-ethylmaleimide-sensitive factor attachment protein receptors) complex. The molecular mechanisms by which Syt1 triggers vesicle fusion remain controversial. This Review introduces and summarizes six molecular models of Syt1: (a) Syt1 triggers SNARE unclamping by displacing complexin, (b) Syt1 clamps SNARE zippering, (c) Syt1 causes membrane curvature, (d) membrane bridging by Syt1, (e) Syt1 is a vesicle-plasma membrane distance regulator, and (f) Syt1 undergoes circular oligomerization. We discuss important conditions to test Syt1 activity *in vitro* and attempt to illustrate the possible roles of Syt1.

Keywords: complexin; neurotransmitter; peptide hormone; SNARE; vesicle fusion; synaptotagmin-1

Neurons and neuroendocrine cells communicate by exocytosis of neurotransmitters at chemical synapses. Neurotransmitters are packaged into specialized organelles called vesicles, i.e., synaptic vesicles and dense core-vesicles. Synaptic vesicles from the presynapse transfer classical neurotransmitters (e.g., glutamate, acetylcholine, gamma-aminobutyric acid, glycine) that regulate electrical signals to postsynapse, whereas dense core-vesicles are responsible for exocytosis of amines, neuropeptides, and hormones, which modulate synaptic activity [1–3].

Dense core-vesicles are minor components (e.g., 1 ~ 2% of density of synaptic vesicles) in the central nervous system such as the neuromuscular junction

[4], thalamus [5], cerebellum [5], and hippocampus [6]. For this reason, chromaffin granules called large dense-core vesicles (LDCVs) have been widely used as a model system to study dense core-vesicles [7]. Despite the differences in the biological functions of synaptic vesicles and LDCVs [2], both types use soluble *N*-ethylmaleimide-sensitive factor attachment protein receptor (SNARE) proteins as the fusion machinery [1–3,8]. Ca^{2+} sensor synaptotagmin-1 (Syt1), which is anchored in vesicle membranes, mediates fast Ca^{2+} -dependent exocytosis of synaptic vesicles in neuromuscular junctions [9] and hippocampal neurons [10], and of LDCVs in chromaffin cells [11,12].

Abbreviations

LDCV, large dense-core vesicle; PI, phosphatidylinositol; PKC, protein kinase C; PS, phosphatidylserine; SNARE, soluble *N*-ethylmaleimide-sensitive factor attachment protein receptors; Syt1, synaptotagmin-1.

Evidence in 1990 showed that the C2 domains of Syt1 have membrane-binding affinity and suggested Syt1 as the Ca^{2+} sensor [13], but further reports have suggested several different molecular mechanisms of how Ca^{2+} triggers Syt1 to initiate vesicle fusion, so the topic has become increasingly enigmatic and even controversial.

This Review focuses on six molecular models to explain the function of Syt1 in Ca^{2+} -dependent vesicle fusion, and discusses the complexity of Syt1 functions: (a) Syt1 triggers SNARE unclamping by displacing complexin, (b) Syt1 clamps SNARE zippering, (c) Syt1 causes membrane curvature, (d) membrane bridging by Syt1, (e) Syt1 is a vesicle-plasma membrane distance regulator, and (f) Syt1 undergoes circular oligomerization.

Syt1 as a Ca^{2+} sensor for fast exocytosis

Neurotransmitters contained in synaptic vesicles are released in response to Ca^{2+} influx at presynaptic nerve terminals [14]. Synaptic transmission at the synapses is fast: the delay from presynapse to postsynapse is around 0.2 ~ 1 ms, depending on temperature; at 37 °C, vesicle fusion occurs within 10 ~ 100 μs after Ca^{2+} influx in the presynapse [15,16]. Syt1 is responsible for fast synchronous exocytosis of synaptic vesicles and LDCV [17,18], but not for asynchronous release, i.e., other Ca^{2+} sensors are involved in asynchronous release (reviewed in [17]). Syt1 that contains two tandem C2 regions, homologous to protein kinase C (PKC) [13] and proposed to be a Ca^{2+} sensor [19], interacts with negatively charged anionic phospholipids in a Ca^{2+} -dependent manner. Knockout of Syt1 in hippocampal neurons causes selective impairment of fast synchronous exocytosis [10]; this result supports the direct evidence that Syt1 is a Ca^{2+} sensor for synchronous release. Mammalian Syt includes 16 isoforms [20], but Syt1, 2, and 9 mediate Ca^{2+} -dependent fusion of synaptic vesicles [21] in neurons, whereas Syt1 and 7 cause Ca^{2+} -dependent LDCV exocytosis [22] in chromaffin cells.

Syt1 is a transmembrane protein in synaptic vesicles and LDCVs. Acidic aspartate residues in two Ca^{2+} -binding C2 domains [13] (C2A and C2B), coordinate three and two Ca^{2+} ions, respectively [23–25]. In addition to these Ca^{2+} -binding loops, the C2B domain contains a polybasic region (KKKK, 324–327; Fig. 1) that is close to Ca^{2+} -binding loops and is enriched with lysine residues [23] that interact with anionic phospholipids [26–32] or the SNARE complex [33–44]. Evidence in 1992 showed that Syt1 interacts with anionic membrane [19] or with SNARE proteins [39] by

electrostatic interaction. Whether Syt1 can interact with anionic phospholipids or the SNARE complex, or both in a physiological ionic environment remains controversial.

Molecular mechanisms of Syt1 to trigger fusion

Model 1: SNARE unclamping by displacing complexin

Complexin I and II are considered to bind to the partially zipper SNARE complex and might block progression of SNARE zippering by replacing synaptobrevin/VAMP-2 binding in the C-terminal part of the SNARE complex [45–48] (Model 1, Fig. 2). However, the hypothesis that complexin clamps SNARE zippering might be applicable only in a specialized synapse and this ‘SNARE clamping by complexin’ hypothesis remains controversial [49], e.g., complexin might promote progression of SNARE zippering [50], complexin might inhibit spontaneous fusion throughout the interaction of membrane [51] and t-SNARE [52], and complexin may clamp vesicle fusion by electrostatic repulsion (reviewed in ref. [49]). Complexin I might block vesicle fusion by the electrostatic repulsion between the vesicle membrane and complexin I in mouse hippocampal neurons [53], but electrostatic repulsion by complexin I showed no significant effect in *C. elegans* neuromuscular junction [54]. The phenotype of complexin I knockout varies among species, so generalization of complexin function is a complicated task. Despite this debate, complexins are generally accepted to be clamping factors that block vesicle fusion in a docking state [45,46,55]. Upon Ca^{2+} influx, Syt1 interacts with the partially zipper SNARE complex and displaces complexin from SNAREs, so the clamping effect of complexin might be released by Ca^{2+} -bound Syt1.

Many groups have demonstrated Syt1 interaction with SNARE proteins [33–44]. The polybasic region in the C2B domain of Syt1 (KKKK, 324–327; Fig. 1) is associated with SNAREs [36–38,56,57]. NMR structure suggested that the polybasic region of the C2B domain (KKKK, 324–327; Fig. 1) interacts electrostatically with acidic residues from syntaxin-1 and SNAP-25 of the SNARE complex, whereas basic residues from Syt1 (bottom end, RR, 398–399; Fig. 1) binds to the membrane [57]; NMR and molecular dynamics simulation suggest that Syt1 and complexin I interact simultaneously with a partially assembled SNARE complex that has distinct binding sites [57]. In contrast, atomic-resolution X-ray crystal structures

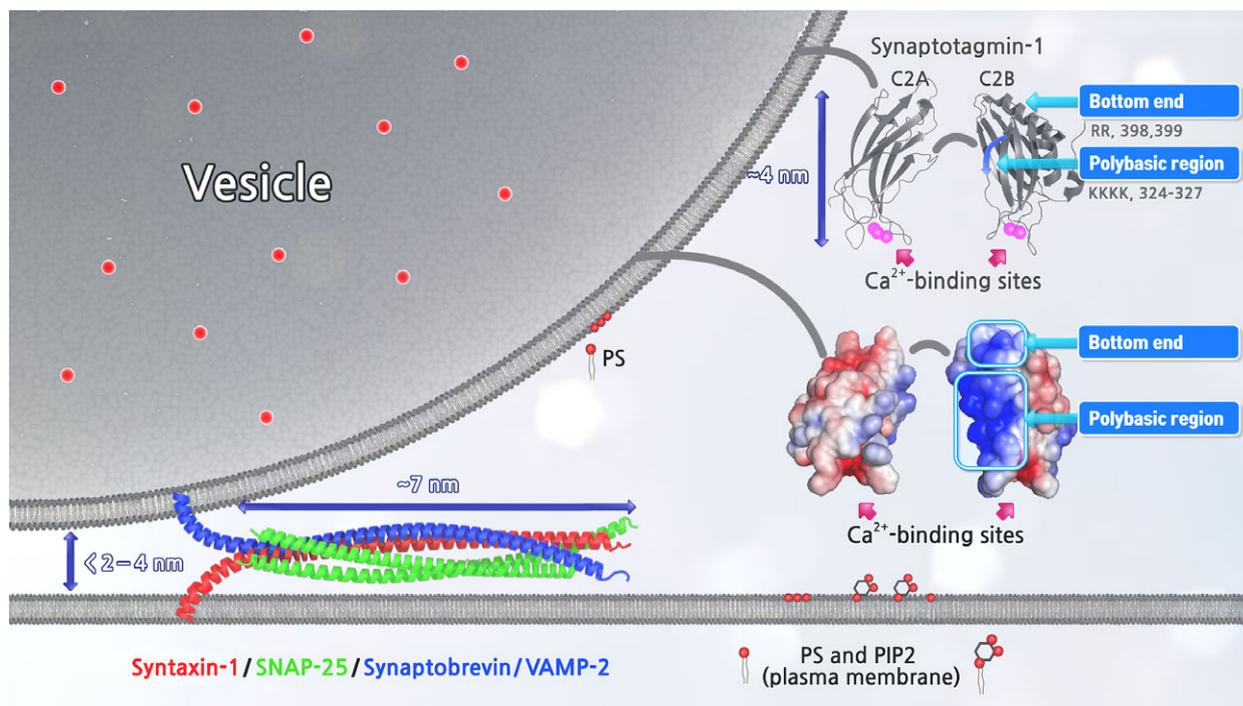


Fig. 1. Structure of the SNARE complex and Syt1. The SNARE motif of the SNARE complex is ~ 7 nm long and synaptic vesicles are tightly docked $< 2 \sim 4$ nm from the plasma membrane; this proximity suggests that SNARE proteins might be partially assembled in a docking state. The C2B domain of Syt1 has a polybasic region with a four-lysine patch (KKKK, 324–327) and bottom end with two arginines (RR, 398–399), which are responsible for the interaction with SNAREs or PIP₂. Protein structures (top) and the electrostatic potential surface (bottom) are visualized using PyMOL Molecular Graphics System (PDB ID: 1BYN for C2A, 1K5W for C2B, and 1SFC for the SNARE complex).

showed that the polybasic region of the C2B domain is not involved in SNARE interaction, but binds to the plasma membrane [58]; and that instead, three different Syt1–SNARE binding sites exist, i.e., the C2A domain of the tertiary interface bridges another SNARE complex and could displace complexin from the SNARE complex [58]. The conserved primary interface involves acidic residues from SNAP-25 and syntaxin-1A and basic residues from Syt1 (bottom end, RR, 398–399; Fig. 1) [58], but an NMR-based model suggests that this region of Syt1 (bottom end, RR, 398–399; Fig. 1) binds to the plasma membrane [57]. These diverse results all indicate that the electrostatic effect dominantly mediates the interaction of Syt1 and the SNARE complex, and that this interaction depends on ionic strength [43,48].

Recent high-resolution crystal structures revealed an additional primed SNARE–Complexin–Syt1 tripartite interface in which two Syt1 molecules simultaneously interact with one SNARE complex on distinct binding sites on primary and tripartite interfaces, termed Syt1–SNARE–Complexin–Syt1 structures [59]. Syt1–SNARE–Complexin–Syt1 complexes might keep the partially zippered SNARE complex in a vesicle-

priming state [59]. Ca²⁺-bound Syt1 causes molecular rearrangements, and ultimately induces full SNARE zippering by liberating or displacing complexin from the Syt1–SNARE–Complexin–Syt1 complex [59].

However, this model in which Syt1 leads to unclamping of SNARE zippering by displacing complexin is still debated. Syt1 binds to SNAP-25 [60–62] and syntaxin-1A [42,62,63] and this interaction is mediated by the electrostatic effect between basic residues of C2AB domain and acidic residues of syntaxin-1A [64] or SNAP-25 [61,65]. Electrostatic interaction between Syt1 and the SNARE complex is mainly observable at low ionic strength [43,48], because under this condition the weak electrostatic interaction strengthens. This interaction of Syt1–SNARE has very weak affinity [31] and is mainly observable at very high concentrations (10–400 μ M) of the C2AB domain of Syt1 [42,60,64–66] and in buffer that has low ionic strength (50–100 mM Na⁺ or K⁺) [43,48,67]. Complexin has a high binding affinity, so it can interact with the SNARE complex independently of ionic strength, but Syt1 fails to bind to the SNARE complex even in the presence of complexin at physiological ionic strength [67].

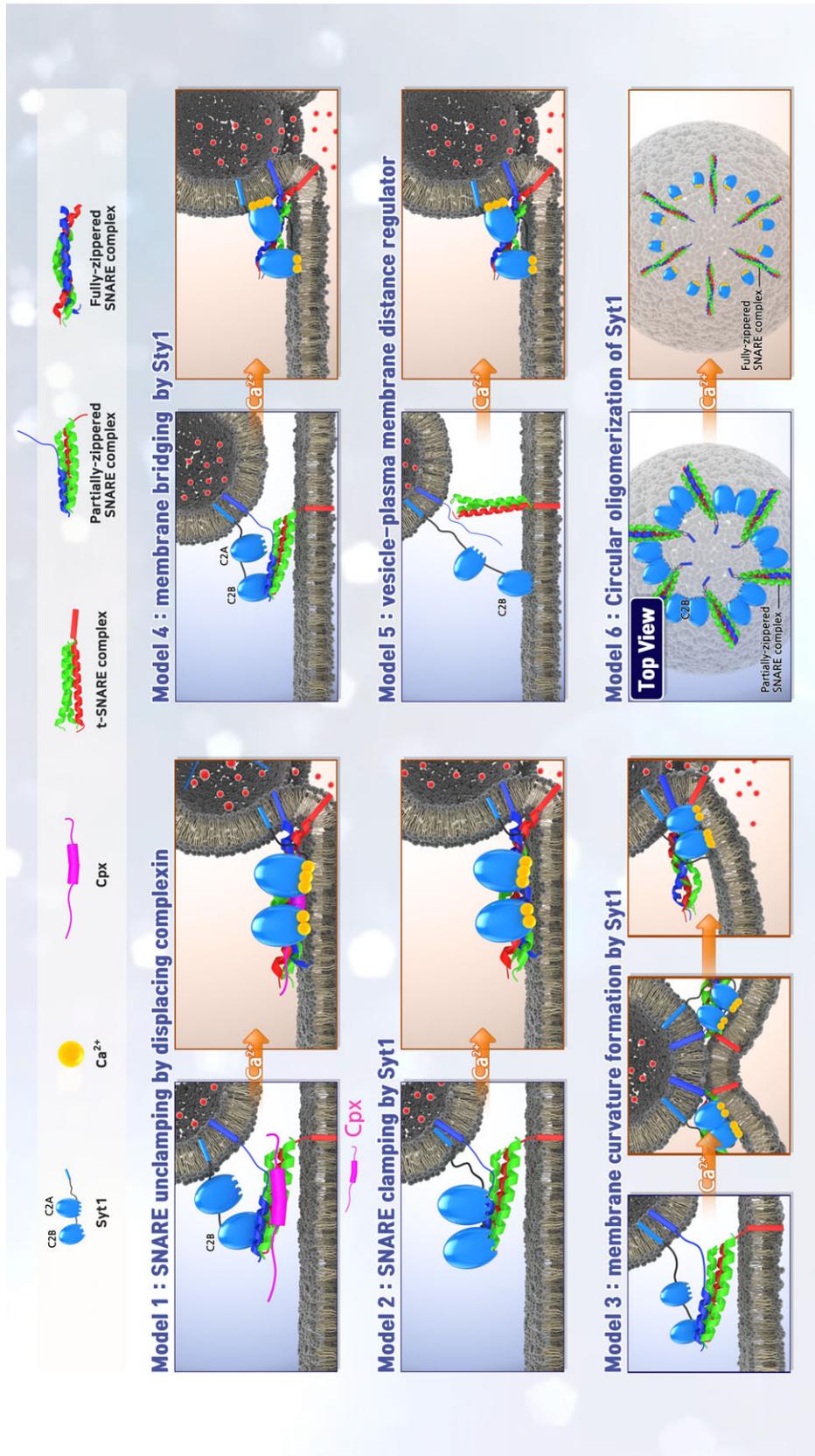


Fig. 2. Six molecular models of Syt1. (Model 1) Syt1-mediated SNARE unclamping by displacing complexin. Complexin clamps the full SNARE zippering before Ca^{2+} arrival. Ca^{2+} -bound Syt1 displaces the complexin so that the partially zippered SNARE complex becomes fully zippered. (Model 2) Syt1-mediated SNARE clamping. Syt1 itself clamps the partially zippered SNARE complex. Ca^{2+} -bound Syt1 is released from clamping the SNARE complex and inserted into the plasma membrane, thus triggering the full SNARE zippering for vesicle fusion. (Model 3) Membrane curvature formation. Ca^{2+} -bound Syt1 forms the local positive curvature in the plasma membrane, thereby lowering energy barrier for SNARE-mediated fusion. (Model 4) Membrane bridging. An equilibrium between partially zippered SNARE complex and repulsion between the two membranes. After Ca^{2+} arrival, C2A and C2B domains bind to opposing membranes to induce the full SNARE zippering and vesicle fusion. (Model 5) Vesicle-plasma membrane distance regulator. The C2B domain of Syt1 binds to the plasma membrane, and after Ca^{2+} arrival, Syt1 bridges two membranes as a 'charge bridge' to initiate the SNARE complex formation to trigger vesicle fusion. (Model 6) Circular oligomerization. A circular oligomerization of Syt1 forms on the plasma membrane by the C2B domains, and thereby blocks full SNARE zippering. Ca^{2+} binding to Syt1 dissociates the ring oligomer so that full SNARE zippering and vesicle fusion occur.

Currently, no direct evidence shows that Syt1 triggers full SNARE zippering by displacing or replacing complexin, which arrests vesicle fusion. Whether complexin clamps SNARE is still under debate, because spontaneous vesicle fusion is either unaffected or reduced in complexin-deficient mammalian neurons (reviewed in Ref. [49]); furthermore, in the physiological ionic environment Syt1 dominantly binds to anionic phospholipids without interacting with the SNARE complex [31,67]. However, complexin could inhibit spontaneous fusion by other processes, for example, the interaction of membrane-complexin [51] and t-SNARE-complexin [52].

Model 2: Syt1 clamps SNARE zippering

Another hypothesis (Model 2, Fig. 2) suggests that Syt1, not complexin, is the clamping factor of SNARE zippering. This hypothesis was proposed because deletion of Syt1 increases spontaneous release (mini-frequency) in many cell types [9,37,68–72], i.e., SNARE clamping inhibits spontaneous release before Ca^{2+} influx. The model proposes that Syt1 clamps SNARE zippering to arrest vesicle fusion before Ca^{2+} triggering, and that Ca^{2+} -bound Syt1 is inserted to the plasma membrane, and thereby causes conformational changes that evoke full SNARE zippering and vesicle fusion.

The C2AB domain of Syt1 has been suggested to be a clamping factor that halts full SNARE zippering by binding to the partially assembled SNARE complex [37]. SNARE clamping by Syt1 may steer the Ca^{2+} -triggered membrane penetration of Syt1 [37]. However, the full-length of Syt1 fails to clamp vesicle fusion, but instead slightly increases the efficiency of Ca^{2+} -independent fusion [29,73–75]; this result strongly argues that Syt1 is not a clamping factor of SNARE assembly. The inhibitory effect of the C2AB domain of Syt1 is also controversial in different systems, e.g., inhibition of Ca^{2+} -independent fusion by the C2AB domain does not occur when purified native vesicles are used [76]. Syt1 inhibits spontaneous release of vesicle fusion, but the mechanisms of SNARE clamping by Syt1 are under debate.

Furthermore, the weak Syt1–SNARE interaction is completely disrupted at physiological ion concentrations that include 150 mM K^+ and 1 mM ATP/ Mg^{2+} [67]; this observation may refute the hypothesis that Syt1 clamps SNARE. However, the interaction of the C2B domain (bottom end, RR, 398–399; Fig. 1) with the SNARE complex is resistant to ionic strength and observable independently of the presence of ATP/ Mg^{2+} ; this result contradicts the

influence of ionic strength on Syt1–SNARE interaction [77]. The electrostatic interaction is tightly coupled to the ionic strength of the buffer [78] and the weak interaction between Syt1 and SNARE proteins is unlikely to be detected in physiological conditions. Ca^{2+} -bound Syt1 has much higher binding affinity to membrane than the SNARE complex, so upon Ca^{2+} triggering, Syt1 only binds to membranes that bear anionic phospholipids [31,67]; this distinction contradicts the physiological role of Syt1–SNARE interaction.

Model 3: Syt1 causes membrane curvature

Ca^{2+} -dependent membrane insertion is the most essential and characteristic property of the C2AB domain of Syt1 [13,19]. Physiological concentration of Ca^{2+} leads to binding of the C2AB domain to negatively charged anionic phospholipids [19,31,42,79–82]. Ca^{2+} -dependent membrane binding of the C2AB domain depends on the concentration of anionic phospholipids, i.e., C2AB domain binding increases with rise in the negative charge density in membranes [83,84]. PIP_2 (–4 net charge at neutral pH) strengthens the binding affinity of Syt1 [31]. Ca^{2+} -binding to aspartate residues in the C2AB domain caused dramatic change in the surface electrostatic potential of the C2AB domain; as a result the positive electrostatic potential of the C2AB domain is attracted to anionic phospholipids, i.e., phosphatidylserine (PS) provides the complete coordination site for Ca^{2+} [25,85], and Ca^{2+} functions as an electrostatic switch that turns off repulsion between anionic phospholipids and acidic residues in the C2AB domain [64]. PIP_2 efficiently potentiates Ca^{2+} -dependent binding of C2AB domain to the target membranes [76,84,86,87] and increases the speed of response of Syt1 insertion by steering the membrane-penetration activity of Syt1 [86]. The polybasic region of the C2B domain (KKKK, 324–327; Fig. 1) increases sensitivity to Ca^{2+} by interacting with PIP_2 in the plasma membrane [26,28–31,86,88–90]. Collectively, the Ca^{2+} -binding loops of Syt1 are inserted to the plasma membrane and the polybasic region of the C2B domain interacts with $\text{PI}(4,5)\text{P}_2$ to synergistically strengthen the binding affinity of Syt1 and increase the dwell time of Syt1 in the plasma membrane [31].

Mutation in the Ca^{2+} -binding sites of Syt1 disrupts synchronous release [83,91], whereas gain-of-function mutation of Syt1 increases Ca^{2+} sensitivity of release [92]; these results agree with *in vitro* data, and support the hypothesis that Ca^{2+} -dependent membrane insertion of Syt1 is critical for Ca^{2+} -dependent vesicle fusion. Additionally, mutations in

the polybasic region of the C2B domain still show Ca^{2+} -dependent exocytosis, but reduced Ca^{2+} -sensitivity of release [88,89].

Two hydrophobic residues in the Ca^{2+} -binding loops of each C2 domain of Syt1 are partially inserted into the inner leaflet of the plasma membrane (~ 10 Å deep) [81]; this action may induce local membrane deformation in the plasma membrane [27,93]. The increased dwell time of Syt1 and deepened insertion of Syt1 into the membrane may destabilize the bilayer and thus contribute to overcome the energy barrier for fusion [31]. The C2B domain of Syt1 causes positive curvature in the plasma membrane in a Ca^{2+} -dependent manner by sequestering PS [94]. Local curvature mediated by membrane insertion of Syt1 may facilitate SNARE-mediated vesicle fusion by lowering the energy barrier of the intermediates including close membrane apposition, fusion stalk formation, and fusion pore openings (Model 3, Figs 2 and 3D) [95–98]. The curvature of plasma membrane might decrease the membrane repulsion force [95,99], and the membrane repulsion becomes minimized by large reduction of the contact area of the two opposing membranes. The energy barrier to stalk formation also decreases because the positive curvature could lower the membrane bending energy required to form a stalk. In addition, the insertion of C2 domains and consequent membrane dimpling increase membrane tension that decreases the energy barrier for both the hemifusion state and fusion pore opening [100].

High concentration of the C2AB domain of Syt1 (~ 10 μM) in the presence of Ca^{2+} tubulates liposomes; this tubulation might result from curvature formation by the C2AB domain [27,93]. However, high concentration of the C2AB fragment is required for tubulation of liposomes. Membrane tubulation by the C2AB domain was only observed by using negative-stain electron microscopy, but not by cryo-electron microscopy [26,101]. Local positive curvature induced by C2AB insertion would facilitate fusion, but local curvature formation is mainly predicted by theoretical simulation [98]; no experiment has yet demonstrated that full-length Syt1 induces endogenous local curvature and membrane protrusion at the area of vesicle docking upon Ca^{2+} triggering. Novel techniques and approaches are expected to show the molecular mechanisms by which Syt1 causes membrane bucking.

Model 4: Syt1 causes membrane bridging

Syt1 cross-links vesicles to the plasma membrane by interactions of the C2A and C2B domain with two different membranes. Before Ca^{2+} arrival, the acidic Ca^{2+} -binding sites of these domains repel membranes,

and this repulsion inhibits membrane fusion [26]. After Ca^{2+} triggering, the C2A and C2B domains bind to opposing membranes, thereby bridging the vesicle and plasma membrane [26,102]. The C2A domain is inserted into the vesicle membrane, but the polybasic region in the C2B domain interacts selectively with PIP_2 , so the C2B domain binds to the plasma membrane [103] (Model 4, Fig. 2).

The linker between the C2A and C2B domain is so flexible that the two C2 domains bridge opposing membrane at ~ 4 nm distance, thereby facilitating vesicle fusion [81,103]. Membrane bridging by Syt1 may be further classified to parallel or antiparallel orientations of two C2 domains, i.e., into an oligomerization model and a direct-bridging model [101]; the linking depends on Syt1 interactions with the vesicle and the plasma membrane [32,73]. Parallel and antiparallel orientations of two C2 domains might have different functions, i.e., parallel orientations of the C2A and C2B domain to the plasma membrane trigger vesicle fusion, whereas antiparallel configuration of domains clamp vesicle fusion [104].

The SNARE complex seems to be partially assembled at a vesicle docking state, and the C2A and C2B domains of Syt1 might face the vesicle and the plasma membrane, respectively. The partial SNARE zipper brings two opposing membrane into close proximity ~ 2 – 4 nm before Ca^{2+} influx [105], then upon Ca^{2+} influx, Syt1 causes membrane bridging up to ~ 4 nm, but the mechanism by which this membrane bridging can lower the energy barrier to fusion is not clear. Membrane bridging by the C2A and C2B domains might decrease the energy barrier for fusion by dehydrating the interbilayer region and stabilizing the primed state (~ 2 nm) (Figs 2 and 3D). However, the physiological ionic environment that includes ATP/ Mg^{2+} disrupts interaction of the C2AB domain with the vesicle membrane [76,106]; these results are evidence that the C2AB domain is only inserted into the PIP_2 -containing plasma membrane to trigger vesicle fusion, not into the vesicle membrane, and may thus refute this model of membrane bridging by the C2AB domains.

Model 5: Syt1 regulates vesicle-plasma membrane distance

The energy barriers for vesicle fusion include dehydration of the phospholipids head groups and steric hindrance; for two opposing membrane to contact, proteins must be moved out of the way. Electrostatic repulsion between the vesicle and plasma membrane might contribute to this energy barrier. The vesicle

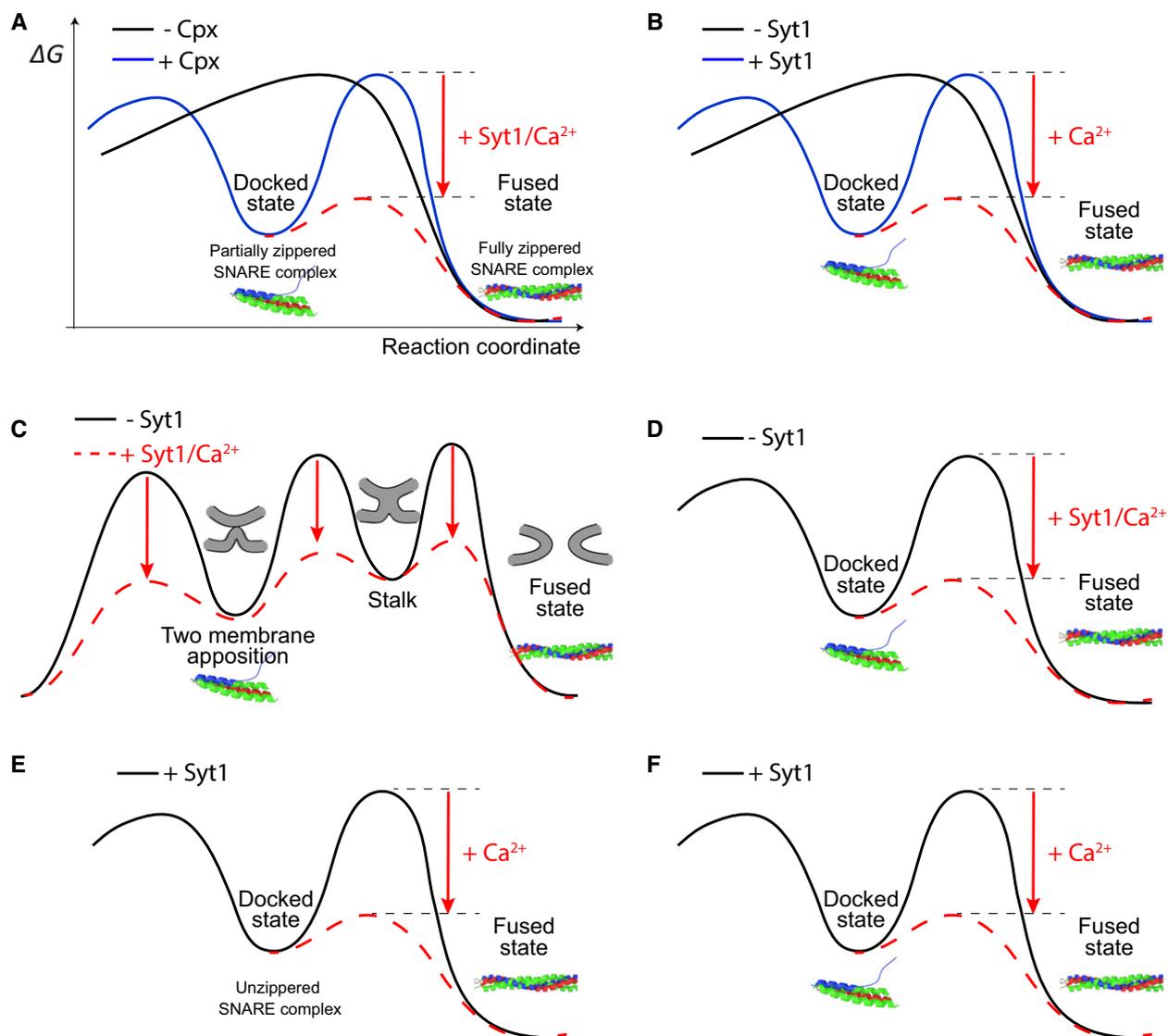


Fig. 3. Energy landscapes of six molecular models of Syt1. (A) Syt1-mediated SNARE unclamping by displacing complexin. Syt1/Ca²⁺ decreases the energy barrier by triggering full SNARE complex formation. (B) Syt1-mediated SNARE clamping. Ca²⁺ binding to Syt1 induces the full SNARE complex formation to decrease the energy barrier for membrane fusion. (C) Membrane curvature formation. Syt1/Ca²⁺ decreases the energy barriers for membrane apposition, stalk formation, and membrane fusion by positive local curvature formation. (D) Membrane bridging. The docked state results from partially zippered SNARE complex formation. Syt1/Ca²⁺ brings the two membranes into close proximity and reduces the energy barrier. (E) Distance regulator. No partial assembly of SNARE proteins at the vesicle docking state. Syt1 controls the distance between the vesicle and plasma membrane and Syt1/Ca²⁺ decreases the energy barrier by bringing two opposing membrane into close proximity. (F) Circular oligomerization of Syt1. C2B oligomerization is released by Ca²⁺ binding to the C2B domain and the dissociation of Syt1 ring decreases the energy barrier for fusion pore opening.

membrane contains the negatively charged phospholipids PS and PI (phosphatidylinositol) [107]; both have -1 net charge, whereas the plasma membrane has PS and PIP₂. Therefore, repulsion between two opposing membrane might block vesicle docking and fusion. This model suggests that SNARE proteins might not be preassembled before Ca²⁺ increase, and that Syt1 contacts the plasma membrane *via* the

polybasic region [75]; this is the major difference from Model 4 (Figs 2 and 3). The polybasic region in the C2B domain is responsible for Ca²⁺-independent binding to PIP₂, and might be involved in vesicle docking [26–32,82,86,88,90]. In a vesicle docking state, electrostatic repulsion expands the distance between two membranes, so SNARE proteins have little chance to meet. Instead, the C2AB domain of Syt1 extends to

interact with the plasma membrane *via* the polybasic region [75]. The polybasic region of the C2B domain tethers two opposing membranes, but electrostatic repulsion prevent SNARE assembly [75] (Model 5, Figs 2 and 3E). Liposomes that contain membrane-anchored Syt1 keep two membranes far ($\sim 7\text{--}8$ nm) apart in the absence of Ca^{2+} [108], and thereby block SNARE zippering.

Upon Ca^{2+} binding, Syt1 functions as a ‘charge bridge’ to bring the two membranes closer together, thus initiating the SNARE nucleation and SNARE-mediated membrane fusion to occur; Syt1 functions as a distance regulator [75]. However, vesicle docking at distance of ~ 8 nm seems unlikely because synaptic vesicles are tightly docked < 2 nm from the active zone and Munc13 and other SNARE regulatory proteins might induce tight vesicle docking *in vivo* [109].

Model 6: Syt1 undergoes circular oligomerization

This model attempts to explain structural studies, and planar ring-shaped Syt1 oligomers that are observed using electron microscopy [106,110,111]. This polymerized oligomerization is conducted by oligomers of the C2B domains. The ring structure is 20–35 nm in diameter, and is composed of 12–20 Syt1 copies [110]. Syt1 rings form on the plasma membrane before Ca^{2+} triggering, and thereby block and clamp the full SNARE zippering and vesicle fusion [106]. Ca^{2+} dissociates the ring oligomerization of Syt1, thus allowing the full SNARE zippering and vesicle fusion [106,110,111] (Figs 2 and 3F).

This buttressed-ring of Syt1 might include the MUN domains of Munc13, i.e., an outer ring composed of six curved Munc13 ‘MUN’ domains surrounds a stable inner ring of Syt1 oligomers, and thereby form a hexameric symmetrical ring structure that clamps SNARE zippering [112]. Circular oligomerization of Syt1 might be consistent with the crystal structure of two Syt1—one SNARE complex [59] in which one C2B domain coordinates the primary site (complexin-independent) and another C2B domain coordinates the tripartite site (complexin-dependent) of the SNARE complex.

Currently, no direct evidence has shown circular oligomerization of Syt1 *in vivo* at the active zone. Visualizing the buttressed rings of Syt1 in presynaptic neurons remains a topic of further study. Although Syt1 mutations that disrupt the ring structure block Syt1 function, the circular oligomerization ring structure of Syt1 is preferentially formed at low ionic strength and reduced at physiological ionic strength [110], i.e., 15 mM KCl for Syt1 ring oligomers [110] and 20 mM KCl for fluorescence correlation

spectroscopy experiments [111]. Nevertheless, this model might describe the geometrical localization between SNARE/Syt1 and the fusion pore.

Discussion

The diversity of Syt1 functions *in vitro* can result from different experimental conditions. Here we describe some experiments to reduce the diversity of Syt1 functions caused by experimental artefacts. 1) *In vitro* fusion data might vary depending on whether full-length Syt1 or the C2AB domain is used for fusion assay. The C2AB domain fails to rescue the phenotype of Syt1 knockout [113] and only full-length Syt1, not the C2AB domain, efficiently reproduces Ca^{2+} -dependent fusion in the *in vitro* reconstitution system [76]. High concentration of the C2AB domain causes clustering and aggregation of liposomes and affect fusion independently of Syt1 activity. 2) Physiological Ca^{2+} concentrations should be used to test Syt1 functions. Ca^{2+} has a biphasic effect to trigger vesicle fusion *in vitro* and in chromaffin cells [67,114], i.e., 10 ~ 100 μM Ca^{2+} increases vesicle fusion, whereas 0.1 ~ 10 mM Ca^{2+} reverses it. Ca^{2+} dose–response curve is an important control experiment to ensure whether or not Ca^{2+} -dependent fusion is mediated by Syt1. 3) A normal physiological ionic environment including ATP/ Mg^{2+} should be used. Syt1 interactions with membrane or SNARE proteins are tightly regulated by the electrostatic effect, and thus Syt1 function and activity vary depending on the ionic strength and salt concentration. Although the effects of low salt condition may be considered because of macromolecular crowding of these proteins [115], Syt1 activity must be confirmed *in vitro* using physiological buffer to reduce the nonspecific effect. SNARE proteins together with SNARE regulatory proteins such as Munc13, Munc18, and even NSF/ α -SNAP create a more realistic reconstitution system to study the Syt1 function (reviewed in [116]).

In a physiological ionic environment, the main binding partner of Syt1 is most likely a membrane that contains anionic phospholipids, rather than the SNARE complex, and the polybasic region of the C2B domain may be responsible for PIP_2 interaction before Ca^{2+} triggering. The possibility still exists that Syt1 interacts with SNARE proteins when vesicles are tightly docked within 0–2 nm of the plasma membrane [109] and the SNARE complex seems to be at least partially assembled. Furthermore, membrane deformation by Ca^{2+} -bound Syt1 might contribute to overcome the energy barrier to fusion. These six models of Syt1 may be compatible and applicable at different

times and in different places, and data obtained using advanced techniques may spawn new models. It is also worth noting that some of the differences between *in vitro* biochemical assays of fusion and *in vivo* neuronal exocytosis may be due indirectly to Syt1 changes in vesicle recycling and cargo sorting, but this Review focuses explicitly on the role of Syt1 in exocytosis.

Conclusions and perspectives

Since the discovery and cloning of Syt1, its mechanisms have been examined intensively. However, molecular models of how Syt1 evokes vesicle fusion remain controversial. Syt1 interacts with SNARE proteins [39] and anionic plasma membrane [19], but the mechanisms have not yet been determined. Major debates are 1) whether Syt1 binds to the SNARE complex or to the membrane or to both [77] in a vesicle docking state in a physiological ionic environment that include ATP/Mg²⁺, 2) how Syt1, together with complexin, controls SNARE clamping to restrict spontaneous fusion before Ca²⁺ rise, 3) whether complexin might be a SNARE clamping factor, and if not, how vesicle fusion stops before Ca²⁺ increase, and 4) how Syt1 unclamps SNARE zippering to trigger Ca²⁺-dependent membrane fusion. We have reviewed six molecular models of Syt1 and discussed the debate about each model. Advances in technologies of cryo-electron microscopy, single-molecule measurements, *in vitro* fusion assay, nanodisc fusion assays for fusion pore nucleation, and force spectroscopy such as optical tweezers, magnetic tweezers and atomic force microscopy will help to increase understanding of the molecular mechanism by which Syt1 triggers vesicle fusion.

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