Mechanism underlying hippocampal long-term potentiation and depression based on competition between endocytosis and exocytosis of AMPA receptors

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N-methyl-D-aspartate (NMDA) receptor-dependent long-term potentiation (LTP) and long-term depression (LTD) of signal transmission form neural circuits and thus are thought to underlie learning and memory. These mechanisms are mediated by AMPA receptor (AMPAR) trafficking in postsynaptic neurons. However, the regulatory mechanism of bidirectional plasticity at excitatory synapses remains unclear. We present a network model of AMPAR trafficking for adult hippocampal pyramidal neurons, which reproduces both LTP and LTD. We show that the induction of both LTP and LTD is regulated by the competition between exocytosis and endocytosis of AMPARs, which are mediated by the calcium-sensors synaptotagmin 1/7 (Syt1/7) and protein interacting with C-kinase 1 (PICK1), respectively. Our result indicates that recycling endosomes containing AMPAR are always ready for Syt1/7-dependent exocytosis of AMPAR at peri-synaptic/synaptic membranes. This is because molecular motor myosin Vb constitutively transports the recycling endosome toward the membrane in a Ca2+-independent manner.

Synaptic plasticity is generally regulated by the release of various neurotransmitters from the presynaptic membrane and/or by varying the density, types, and properties of neurotransmitter receptors at the postsynaptic membrane. NMDA (N-methyl-D-aspartate) receptor-dependent LTP and LTD of signal transmission in excitatory neurons, such as hippocampal pyramidal neurons, is thought to underlie the formation of neuronal circuits during learning and memory1–3. Fast excitatory neurotransmission in the mammalian brain is predominantly mediated by the AMPA (α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid) receptor (AMPAR) at the postsynaptic membrane. AMPAR is tetrameric ion channel composed of the subunits GluA1–A4 (or named as GluR1–R4). In hippocampal pyramidal neurons, the GluA1/A2 heterotetramer (Fig. 1a) is the most dominant AMPAR subtype, followed by the GluA2/A3 heterotetramer4. It is well established that AMPAR trafficking in postsynaptic neurons plays a decisive role in the induction of LTP and LTD5–8, whereas the dominant pathway and regulatory mechanism for AMPAR trafficking remains a controversial issue.

NMDA receptor-dependent LTP and LTD are triggered by standard high-frequency stimulations (e.g., one or more trains of 100 Hz stimulation)9,10 and low-frequency stimulations (LFS; e.g., 700–900 pulses at 1 Hz)11–14, respectively. It is widely accepted that a high rise and lower rise in intracellular Ca2+ concentrations mediated by NMDA receptor activation are required for the expression of LTP and LTD, respectively. Nevertheless, it
has been reported that a channel blocker for Ca\(^{2+}\) influx through the NMDA receptor does not block LTD in the hippocampal neurons of immature rodents, thereby suggesting that LTD is attributed to an ion-channel-independent, metabotropic form of NMDA-receptor signaling\(^{15}\). Subsequently, it was carefully reexamined how varying extracellular Ca\(^{2+}\) levels and blocking NMDA-receptor channel ion flux with the same channel blocker affected the induction of LTD. It was reconfirmed that the LTD induced by a standard LFS in hippocampal neurons of both adult and immature rodents was dependent on ionotropic NMDA-receptor signaling\(^{16}\).

For more than 20 years, it has been known that Ca\(^{2+}/\text{calmodulin-dependent protein kinase (CaMKII)}\) activity is necessary for LTP; Ca\(^{2+}\) ions incorporate into the cell through NMDA receptor and bind to calmodulin (CaM). Consequently Ca\(^{2+}/\text{CaM}\) is bound to CaMKII. If CaMKII is knocked out, Ca\(^{2+}/\text{CaM}\) binds directly to NMDA receptors, resulting in inactivation of the NMDA receptor channel and an accordingly significantly lower rise in Ca\(^{2+}\) concentrations\(^{17,18}\). On the other hand, if Ca\(^{2+}/\text{CaM}\)-bound CaMKII binds to the NMDA receptor and inhibits the direct binding of Ca\(^{2+}/\text{CaM}\) to the NMDA receptor, this maintains the channel activity of the NMDA
receptor. This is one of the most important CaMKII functions at the early stage of NMDA receptor-dependent LTP induction process. In addition, the Ca\(^{2+}\)/CaM-CaMKII bound to the NMDA receptor phosphorylates GluA1\(^{19}\), resulting in increased channel conductance of AMPAR\(^{20}\). The activation of AMPAR by CaMKII also plays a crucial role in the induction of LTP. In the present study, we build a network model to reproduce NMDA-receptor dependent bidirectional synaptic plasticity of hippocampal neurons. All of the molecular mechanisms incorporated into the model work downstream of these CaMKII functions; thus, the normal activities of CaMKII are necessary and implicitly assumed in our network model. In fact, NMDA receptor-dependent Ca\(^{2+}\)- influx is introduced in the network model as the input for LTP and LTD simulations. In addition, it is assumed that the strength of LTP and LTD induction is in proportion to the change in the AMPAR population on the postsynaptic membrane. Thus, the effects of GluA2-lacking Ca\(^{2+}\)-permeable AMPARs are not taken into consideration in our simulations. However, this assumption does not influence the prediction of LTP expression by our network model, because it has been demonstrated that LTP in the hippocampal CA1 region does not require insertion or activation of GluA2-lacking AMPARs\(^{21}\).

NMDA-receptor dependent LTP is thought to occur by incorporating AMPARs into the synaptic membrane through either the exocytosis of the recycling endosome at peri-synaptic/synaptic membranes\(^{22,25}\), the cell surface long-range lateral diffusion of AMPAR from the dendrites/extrasynaptic region\(^{26-28}\), or a combination of the two. Recently, Penn et al. examined the effects of crosslinking immobilization of pre-existing membrane AMPARs on early LTP and observed slow continuous development of early LTP without short-term potentiation (STP)\(^{29}\). They also observed that blocking exocytosis of AMPARs completely abolished early LTP and caused only STP induction\(^{26}\). The former and latter respectively indicate the following: (1) lateral diffusion movement of AMPARs exocytosed at the synaptic/peri-synaptic membranes as well as the extrasynaptic membranes is significantly suppressed due to obstacle effects arising from AMPAR crowding, as predicted by computer simulations\(^{30}\); (2) although the recruitment of pre-existing membrane AMPARs from the peri-synaptic membrane works for the induction of STP, it is insufficient for maintaining the expression of early LTP. Therefore, the diffusion dynamics on the membrane strongly affect the form of LTP expression and play a crucial role in the rapid short-range diffusion relocation of AMPARs from the peri-synaptic to the synaptic membrane. It was also demonstrated by the duration of STP\(^{30}\) that the short-range diffusion relocation takes, at most, only 2 min. On the other hand, the observed delay of LTP induction caused by the obstacle effects provides no definitive evidence that the long-range lateral diffusion pathway of AMPARs from the dendrites/extrasynaptic region is the predominant pathway for early LTP than the short-range diffusional relocation pathway of exocytic AMPARs from the peri-synaptic membranes. Indeed, exocytosis of GluA1, not only in dendrites but also in synaptic spines including synaptic/peri-synaptic membranes, has been demonstrated using the SEP-GluA1 imaging technique, which resolved the controversy\(^{27,31}\). Alternatively, that observation does not necessarily exclude the short-range diffusional relocation pathway from the promising AMPAR recruitment pathway during LTP. This is because the impaired diffusion movement of AMPARs exocytosed at the peri-synaptic membranes, which is caused by the obstacle effects due to jamming/crowding of crosslinked AMPARs, can yield a delay to early LTP. This hypothesis suggests that the total amount of AMPARs exocytosed at the peri-synaptic membranes should be sufficiently greater than that at the synaptic membrane due to the larger area of peri-synaptic membranes.

In general, the normal lateral diffusion process is isotropic; thus, the flux of lateral diffusion from a place in the extrasynaptic region toward the synaptic membrane and from this place in the opposing direction are in equilibrium (for instance\(^{25}\)). Therefore, the cell surface lateral diffusion movement is thought to be ineffective for long-range directional transport of AMPARs. Furthermore, the Ca\(^{2+}\)-dependent/independent biased surface diffusion mechanism of AMPAR toward the synaptic membrane from the dendrites/extrasynaptic region remains unclear. The long-range transport of AMPARs should necessitate directional non-equilibrium active movement, e.g., driven by molecular motors that consume ATP as fuel (for instance\(^{35}\)).

**Method**

**Network model.** In the network model presented here, we incorporate elemental processes involved in AMPAR trafficking cycles that have been characterized experimentally for hippocampal postsynaptic neurons (Fig. 1). The AMPAR trafficking cycle is constructed around the phosphorylation/dephosphorylation dynamics of AMPAR at the synaptic membrane (Fig. 1c,d)\(^{34-36}\). The endocytosis of vesicles containing AMPAR (Fig. 1e,f)\(^{37,38}\), the transport of the recycling endosomes\(^{39,40}\) by molecular motor myosin \(V_s\)\(^{10-12}\) (Fig. 1g), and exocytosis of the recycling endosomes\(^{22,23}\), followed by the incorporation of AMPARs into the peri-synaptic/synaptic membranes (Fig. 1h). The key facilitators of the AMPAR trafficking cycle\(^{41}\) are summarized as follows: the GluA1/A2 heterotramer (Fig. 1a); A-kinase anchoring protein 150 (AKAP150)\(^{43,44,45}\), which dimerizes (as for long-range directional transport of AMPARs. Furthermore, the Ca\(^{2+}\)-dependent/independent biased surface diffusion mechanism of AMPAR toward the synaptic membrane from the dendrites/extrasynaptic region remains unclear. The long-range transport of AMPARs should necessitate directional non-equilibrium active movement, e.g., driven by molecular motors that consume ATP as fuel (for instance\(^{35}\)).

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NMDA receptors induces the active state of the Ca\(^{2+}\)-sensor PICK1 and then the endocytosis of AMPARs, which are comprised of two GluA1s with dephosphorylated S845 and two GluA2s with phosphorylated S880\(^{37,38,47,55}\). Consequently, the endocytic vesicles containing these AMPARs undergo diffusion in the cytosol (Fig. 1f). At this point, it is assumed that the S880 of GluA2s in recycling endosomes are dephosphorylated by PP2A\(^{38,47}\), PICK1 dissociates from GluA2, whilst GRIP1 binds to GluA2 (Fig. 1g)\(^{43,46}\). It has been demonstrated using GRIP1 and/or GRIP2 knock-out mice that GRIP1 and GRIP2 regulate AMPAR trafficking\(^{46}\). However, for simplification, we assume that the functions of GRIP2 are also taken into consideration through the model of GRIP1 in the present study.

We explicitly take into consideration the active transport of recycling endosomes containing AMPARs by myosin V\(_b\) in our network model. Myosin V\(_b\) binds to the recycling endosome via Rab11 and transports it toward the peri-synaptic/synaptic membrane\(^{31,40–42}\) (Fig. 1g). Although Ca\(^{2+}\)-dependent activity of myosin V has been reported\(^{38}\), the run speed and mean run length are almost constant at the Ca\(^{2+}\) concentrations that we consider in this study\(^{59}\). Therefore, we model the myosin-V\(_b\) transport of recycling endosomes as a Ca\(^{2+}\)-independent constitutive movement toward the peri-synaptic/synaptic membrane, which is driven by ATP hydrolysis energy. Exocytosis of the recycling endosomes transported by myosin V\(_b\) is mediated at the peri-synaptic/synaptic membrane by the Ca\(^{2+}\)-sensor synaptic vesicle protein synaptotagmin 1 (Syt1), together with synaptotagmin 7 (Syt7) (not shown in Fig. 1), synaptobrevin-2/VAMP2, and complexin (not shown in Fig. 1), amongst others\(^{22,23,49,50,60,61}\). As a result, the AMPARs are incorporated into the peri-synaptic/synaptic membrane (Fig. 1h). In fact, exocytosis of GluA1 not only in dendrites but also in synaptic spines including peri-synaptic/synaptic membranes has been demonstrated using the SEP-GluA1 imaging technique\(^{35}\). The localization of Syt1 at the peri-synaptic/synaptic membranes, which has been observed for hippocampal postsynaptic neurons\(^{40}\), also supports the Syt1-mediated exocytosis pathway of the recycling endosomes. The exocytic AMPARs incorporated into the peri-synaptic/synaptic membrane are relocated into the synaptic membrane via a local diffusional movement, which takes, at most, only 2 min, as shown by the observation on the duration of STP\(^{26}\). In the present study, we did not introduce enough fine space resolution into our model network to identify whether AMPARs are located at the peri-synaptic or synaptic membrane. However, the ~2-min delay of AMPAR incorporation into the synaptic membrane after exocytosis at the peri-synaptic membrane would be taken into consideration via effective rates for synaptotagmin-dependent exocytosis. In addition, the diffusional relocation dynamics of AMPARs at the synaptic membrane plays a crucial role in meeting and interacting with the signaling complex AKAP150, thereby resulting in the stabilization of AMPARs at the synaptic membrane by tethering to AKAP150 via SAP97 and GRIP1, as mentioned above.

As a result, we propose the network model for bidirectional hippocampal synaptic plasticity that consists of 285 reaction equations for 191 components, including: (1) Ca\(^{2+}\) dynamics, (2) the phosphorylation/dephosphorylation dynamics of the tetrameric AMPAR ion channel subtype GluA1/A2, (3) the endocytosis/exocytosis dynamics of the AMPARs, mediated respectively by the Ca\(^{2+}\)-sensors PICK1 and Syt1, and (4) the recycling endosome active transport by molecular motor myosin V\(_b\) toward the peri-synaptic/synaptic membrane (Fig. 2, Supplementary Table S2). Thus, our network model involves the previously proposed model on bidirectional synaptic plasticity based on the phosphorylation/dephosphorylation of GluA1 S845\(^{35}\). The validity of the network model is also demonstrated by reproducing the impairments of LTP and LTD caused respectively by genetic chemical inhibition of myosin V\(_b\) transport\(^{36}\) and by AKAP150\(_{\pm}\)PIX knock-in mice, which selectively disrupt the anchoring of PP2B to AKAP150\(^{50}\). The network model reveals that the competition between exocytosis caused by Syt1, and endocytosis caused by PICK1 depends on transient increases in intracellular Ca\(^{2+}\) concentrations, which therefore regulate AMPAR trafficking resulting in LTP or LTD. The obtained results also indicate that the constitutive active transport of the recycling endosomes by myosin V\(_b\) increases the basal concentration of the recycling endosomes localized on the peri-synaptic/synaptic membrane surface, so that the AMPARs, which are ready for Syt1-mediated exocytosis, can be immediately incorporated into the postsynaptic membranes by LTP stimulation. This insertion mechanism resolves the long-standing contradiction between the prompt LTP induction and the several-minute delay on the starting time of myosin V\(_b\) transport following LTP\(^{12}\).

**Simulations.** We built the network model for hippocampal LTP and LTD which is consisted of a well-mixed single compartment. The ordinary differential equations for the network model comprised of 285 reaction equations on 191 components were solved using COPASI biochemical system simulator (ver. 4.23)\(^{53}\). The detailed description of the model including components, reactions, and parameters can be found in the SI text. We employed the steady-state concentrations as the initial condition and performed the simulations of the LTP and LTD inductions for 5,400 s.

**Results**

**Simulated Ca\(^{2+}\) pulses induce LTP and LTD.** The magnitudes of LTP and LTD depend on experimental protocols, though typical forms of LTP and LTD have been found. For instance, it has been observed that high-frequency stimulation (HFS) at 100 Hz for 1 s sharply induces LTP, increasing synaptic transmission up to ~200% compared to basal levels\(^{37,38,64}\). The magnitude drops rapidly to ~150%, and then slowly decreases toward basal transmission level. Our network model reproduces the change in the membrane AMPAR level that is qualitatively consistent with experimentally observed LTP induction (Fig. 3a), when we mimic a rapid transient increase in intracellular Ca\(^{2+}\) concentration caused by HFS-induced Ca\(^{2+}\) influx through NMDA receptors, which is the input for LTP simulation, using a single Gaussian function (Fig. 3b). On the other hand, it has been found experimentally that induction of LTD requires ~2,000 Hz (700–900 pulses at 1 Hz)\(^{11,13}\). Synaptic transmission gradually decreases down to ~60%\(^{11,13}\), and then slowly increases toward the basal transmission level\(^{11,14}\). When the transient rise of intracellular Ca\(^{2+}\) caused by LFS is modelled using three Gaussian functions (Fig. 3b), our
network model reproduces the experimentally observed membrane AMPAR level for LTD\(^{31}\) (Fig. 3a). In addition, we found that the network model reproduces LTP and LTD inductions under a wide range of Ca\(^{2+}\) peak pulse amplitudes (see Supplementary Fig. S1). Furthermore, we confirmed that these LTP and LTD inductions were held even if we used two sigmoid functions to mimic Ca\(^{2+}\) influx instead of the Gaussian functions (see Supplementary Fig. S2). These results indicate the validity of the network model for bidirectional synaptic plasticity. Here, it is noted that the network model reproduces the slow reduction in the amplitude of LTP and LTD toward the basal condition, which has been observed experimentally. Thus, the LTP yielded by the network model, which is based on AMPAR trafficking, corresponds to the early phases of LTP. The relaxation of LTP and LTD is mediated by constitutive endocytosis and exocytosis of AMPAR, thus indicating that the constitutive flux of the recycling endosome occurs under basal conditions.

**PICK1** is activated by both LTP and LTD stimulation whereas synaptotagmin 1 is mainly activated by LTP stimulation. In our network model, the Ca\(^{2+}\)-sensors Syt1 and PICK1 play a dominant role in the exocytosis and endocytosis of AMPARs that regulate AMPAR trafficking during LTP and LTD induction. We built biochemical models of Syt1 and PICK1 based on experimentally determined association constants of Ca\(^{2+}\) ions to Syt1\(^{60}\) and PICK1\(^{55}\) (see the SI text). Syt1 has two Ca\(^{2+}\)-binding domains, C2A and C2B, which bind three and two Ca\(^{2+}\) ions and directly interact with the recycling endosome and plasma membrane, respectively\(^{60}\). PICK1 has two Ca\(^{2+}\)-binding domains, one each on the N-terminus and C-terminus, which interact with the phosphorylated S880 of GluA2 and the synaptic plasma membrane, respectively\(^{37,38}\). It has been demonstrated that Ca\(^{2+}\)-binding site mutations of Syt1 in both the C2A and C2B domains block hippocampal
therefore, it would be appropriate to use Ca²⁺-binding constants of Syt1 to model a Ca²⁺-dependent regulating factor on the exocytosis mediated by Syt1 together with Syt7, synaptobrevin-2/VAMP2, and complexin, amongst others. During LTP stimulation, the concentrations of multiple Ca²⁺-binding species of Syt1, \((\text{Ca}^2+)_{\text{ASyt1}}\), \((\text{Ca}^2+)_{\text{ASyt1}}(\text{Ca})_{\text{BSyt1}}\), and \((\text{Ca}^2+)_{\text{ASyt1}}(\text{Ca})_{\text{BSyt1}}\), rapidly rise in relation to the increase in \([\text{Ca}^2+]_{\text{ASyt1}}\). Here and hereafter, we use square brackets to refer to concentrations. However, at this scale, \([\text{Ca}^2+]_{\text{ASyt1}}(\text{Ca})_{\text{BSyt1}}\) does not show a significant increase (Fig. 3d). Therefore, both Syt1 and PICK1 are activated by LTP stimulation. Nevertheless, LTP is induced because the Syt1-mediated exocytosis overcomes the PICK1-mediated endocytosis. In contrast, at this scale during the LTD stimulation, increases in the concentration of multiple Ca²⁺-binding species of Syt1 are not seen (Fig. 3f), whereas \([\text{Ca}^2+]_{\text{PICK1}}\) and \([\text{Ca}^2+]_{\text{PICK1}}\) rise in relation to the rise in \([\text{Ca}^2+]\) (Fig. 3e). As a result, the PICK1-mediated endocytosis overcomes the Syt1-mediated exocytosis, causing the induction of LTD (the numerical results will be provided below). The difference in Ca²⁺-dependent activity of PICK1 and Syt1 during the LTP and LTD stimulation is attributable to that in Ca²⁺-binding affinity of these proteins (see Supplementary Fig. S3). Graupner and Brunel proposed that the total times when Ca²⁺ transient spends above depression and potentiation thresholds of synaptic transmission determined LTD and LTP induction²⁵. The former and latter

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**Figure 3. Hippocampal LTP and LTD are regulated by the activation of Ca²⁺-sensors Syt1 and PICK1 in response to Ca²⁺ influx.** (a) Time course of the membrane AMPAR population, indicating induction of LTP and LTD. Here, 100% represents the basal AMPAR population at the membrane. (b) Ca²⁺-pulse concentrations corresponding to the LTP and LTD stimulation are shown on the left and right axis, respectively. (c-f) The concentrations of Ca²⁺-binding species of PICK1 (c,e) and Syt1 (d,f) as a function of time (t) during LTP (c and d) and LTD induction (e,f). In (f), the concentration of multiple Ca²⁺-binding species other than \((\text{Ca}^2+)_{\text{ASyt1}}(\text{Ca})_{\text{BSyt1}}\) is too small to see at this scale, indicating that Syt1 is mostly not activated. Therefore, Ca²⁺-dependent exocytosis mediated by Syt1 occurs during LTD.
might correspond to Ca^{2+} concentrations that activate PICK1-dependent endocytosis and Syt1-dependent exocytosis in our network model.

It has been reported that the regulatory mechanism of AMPAR trafficking on hippocampal synaptic plasticity of rodents is development dependent. In fact, LTD is modestly affected in juvenile PICK1-knock out (KO) mice, whereas LTD induced by LFS is obviously reduced in PICK1-KO adult mice\textsuperscript{66}. Although many factors other than PICK1 work in Ca^{2+}-dependent endocytosis of AMPAR during LTD especially in hippocampal neurons of juvenile rodents, PICK1 plays a dominant role in the regulation of Ca^{2+}-dependent endocytosis in adult rodents together with other proteins including clathrin and dynamin\textsuperscript{67}. Thus, the regulatory mechanism of hippocampal bidirectional synaptic plasticity presented in this study is valid, especially for adult rodents. Very recently, it was demonstrated using synaptotagmin 3 (Syt3) KO that Syt3 is actually involved in the endocytosis of AMPARs during LTD\textsuperscript{68}. However, the Ca^{2+} affinity of Syt3 has been reported as being tenfold higher than that of Syt1\textsuperscript{69}, at levels more similar to PICK1. Therefore, Syt3-mediated endocytosis does not prevent the Ca^{2+}-dependent regulatory mechanisms of exocytosis and endocytosis presented here, and the network model would reproduce the bidirectional synaptic plasticity even if it was also taken into consideration.

Figure 4. Competition between exocytosis and endocytosis of AMPARs yields LTP and LTD. (a,c) Total excess fluxes of exocytosis, endocytosis, and myosin Vb transport of recycling endosomes during (a) LTP stimulation and (c) LTD stimulation. (b,d) The time course of concentrations for predominant components during (b) LTP and (d) LTD.
...tions and our simulation results showing an impairment of both the LTP and LTD (see Supplementary Fig. S4) are the convincing evidence that supports our competition mechanism of bidirectional synaptic plasticity.

**Endocytic vesicles newly generated during LTP and LTD are transported by myosin Vb toward the peri-synaptic membrane.** Endocytic vesicles containing AMPARs newly generated during the induction of LTP and LTD undergo diffusion in the cytosol as a recycling endosome. During the diffusion movement these recycling endosomes bind to molecular motor myosin Vb via Rab11 and are actively transported by it toward the peri-synaptic/synaptic membrane. The long-term flux by myosin Vb transport is found in Fig. 4a,c. This indicates that the myosin Vb transport of recycling endosomes continues for a while after LTP and LTD are induced. Indeed, such active transport continues for ~7 and ~13 min from the onset of the LTP and LTD stimulation, respectively (Fig. 4a,c). The result obtained for LTP is consistent with experimental observations where the cooperative movements of recycling endosomes and myosin Vb molecules start a few minutes later than the LTP induction, and continue for several minutes.

Recycling endosomes transported by myosin Vb are localized on the surface of the peri-synaptic membrane, and thus are already prepared for exocytosis. We assume that the myosin Vb active transport takes place in a Ca^{2+}-independent manner based on experimental observations. Thus, the recycling endosomes in the cytosol are constitutively transported by myosin Vb toward the peri-synaptic and synaptic membranes. Consequently, the recycling endosomes are expected to become localized on the membrane surface under basal conditions. In fact, we see that the most dominant components in the cytosol under basal conditions (t = 0 s) are pre-exocytic recycling endosomes bound to the peri-synaptic/synaptic membrane surface, namely, R1R2psd(GRIP1),Myo5VSYt1 (Fig. 4b,d). Such localization is necessary for the immediate exocytosis mediated by Syt1 after LTP stimulation. Specifically, this causes the rapid incorporation of AMPARs into the peri-synaptic/synaptic membranes during LTP induction. Sharp decreases in [R1R2psd(GRIP1),Myo5VSYt1] occur immediately after LTP stimulation until it is depleted by the Syt1-mediated exocytosis. Simultaneously, the AMPAR population at the membranes increases due to Syt1-mediated exocytosis, then immediately begins to decrease after it reaches its peak (see Fig. 3a). Such a rapid decrease in the AMPAR population following the prompt increase is interpreted as being due to PICK1-mediated endocytosis following Syt1-mediated exocytosis, as seen in the excess fluxes during LTP (Fig. 4a). The increase in [R1R2psd(GRIP1),Myo5VSYt1] following depletion (Fig. 4b) is therefore attributed to the myosin Vb-transport of the endosomes newly internalized into the cytosol through endocytosis (Fig. 4a,c). Likewise, endocytic vesicles internalized during LTD induction are also transported by myosin Vb and are present in greater quantities on the membrane surface than under basal conditions (Fig. 4d), thus playing an important role in exocytosis due to subsequent LTP stimulation.

**Discussion**

**Assessment of the validity of the network model for hippocampal LTP and LTD.** To confirm the validity of the network model for bidirectional synaptic plasticity, we applied the model to two experimental observations: (1) reduction in LTP induction through genetic chemical inhibition of myosin Vb transport, and (2) impaired LTD induction in AKAP150ΔPIX knock-in mice where the anchoring of PP2B to AKAP150 is selectively disrupted. The former is useful to reveal how myosin Vb transport affects LTP induction. This is not a trivial issue, because the myosin Vb transport starts a few minutes later than the LTP induction, and the movement of recycling endosomes continues for ~10 min. The latter is important to reveal how the competition between the phosphorylation/dephosphorylation of GluA1 S845 by PKA and PP2B affects LTD induction. Indeed, on the basis of localization of AMPARs at the synaptic membrane regulated by phosphorylation of GluA1 S845, a model of bidirectional synaptic plasticity has been proposed.

Our network model reproduces the experimentally observed reduction in LTP due to the inhibition of myosin Vb transport (Fig. 5a), i.e., that the LTP magnitude is lower in the early stage (~ t = 700 s) than in the wild-type model, and that the following reduction of LTP toward basal levels is accelerated. The model also reproduces experimentally observed LTD impairment by disrupting dephosphorylation of GluA1 S845 by PP2B (Fig. 6a). Additional demonstrations of the network model are presented in Supplementary Fig. S5. These simulation results are also useful to further support the validity of the bidirectional regulatory mechanism of the network model on the induction of both LTP and LTD.

**The accelerated reduction of LTP toward basal levels reflects inhibition of myosin Vb transport.** Inhibition of myosin Vb transport does not affect the rapid increase in the membrane AMPAR population just after the onset of LTP stimulation (Fig. 5a). However, following this rapid increase, the AMPAR population steeply drops down to lower levels than that of the wild-type model (~ t = 700 s), showing an impairment of LTP induction. The steep decrease after the initial increase can be attributed to PICK1-mediated endocytosis, which occurs concurrently during LTP induction (Fig. 4a). Normalized endocytic fluxes, defined as the total endocytic flux divided by the basal AMPAR, indicate that the ratio of endocytic AMPARs relative to the total amount of membrane AMPARs is increased by the inhibition of myosin Vb transport (Fig. 5b). Whilst the increase in the normalized endocytic flux is unexpected, it can be interpreted through changes in the population that are related to PICK1-bound species, to which myosin Vb does not directly bind (Supplementary Fig. S6). These observations show that the reduction in LTP in the early stage (~ t = 700 s) is attributable to indirect effects of the inhibition of myosin Vb transport. In fact, the cooperative movement of recycling endosomes and myosin Vb molecules starts several minutes later than LTP induction.

On the other hand, the accelerated reduction in LTP magnitude toward basal levels compared with the wild-type model can be interpreted as the direct effects due to inhibition of myosin Vb transport. Indeed, the...
concentration of recycling endosomes to which myosin Vb binds, \([R1R2_{endo(GRIP2)}\text{MyoV}]\), is significantly increased due to the inhibition of myosin Vb transport not only under basal conditions \((t = 0 \text{ s})\) but also at times after the induction of LTP (e.g., \(t = 1,000 \text{ s}\), Fig. 5d). In contrast, in the wild-type model, \([R1R2_{endo(GRIP2)}\text{MyoV}]\) slightly increases only during LTP stimulation, and immediately vanishes due to myosin Vb transport (Fig. 5c). The slow increase in the concentration of the recycling endosomes on the membrane surface, \([R1R2_{psd(GRIP2)}\text{MyoV}]\), by inhibiting myosin Vb transport reflects the rate-limiting process in the constitutive cycle of AMPAR trafficking, which results in a faster reduction in LTP toward basal levels42.

**Phosphorylation/dephosphorylation dynamics of AMPARs at the synaptic membrane regulate AMPAR trafficking.** AKAP15034,44,45 forms a signaling complex with PKA, PKC, and PP2B (Calcineurin, CaN, Fig. 1b) and affects the phosphorylation/dephosphorylation dynamics of the tetrameric AMPAR ion channel GluA1/A2 (Fig. 1c,d). The S845 site of GluA1 is phosphorylated and dephosphorylated by PKA and PP2B, respectively. The enzymes PKA and PP2B control the localization of GluA1/A2 to the synaptic membrane by tethering GluA1 to AKAP150 via SAP97, due to the fact that SAP97 preferentially binds to the phosphorylated S845 site of GluA135,36. However, it remains unclear how these opposing effects of PKA and PP2B are regulated during the induction of LTD. To address this issue, AKAP150ΔPIX knock-in mice, where the anchoring of PP2B to AKAP150 is disrupted, have been developed. Using the knock-in mice, it has been demonstrated that inhibition of the interaction between AKAP150 and PP2B impairs hippocampal LTD induction36. In this study, we modeled hippocampal synapses for AKAP150ΔPIX knock-in mice by suppressing the dephosphorylation rate of GluA1 S845 by PP2B (Supplementary Table S5). The LTD induction is impaired in the AKAP150ΔPIX model compared with the wild-type model (Fig. 6a). In Fig. 6c,d, R1R2(pS2) species that bind SAP97, R1(pS-SAP)R2(pS2), and R1(pS-SAP)R2(pS), show larger populations than species that do not bind SAP97 by disrupting PP2B-anchoring to AKAP150. This is because dephosphorylation of GluA1 S845 is suppressed in the AKAP150ΔPIX model. Consequently, the population of the main species in preparation for endocytosis, such as R1R2(pS2-PICK1) and R1R2(pS-CaPICK1), is reduced in the AKAP150ΔPIX model, resulting in a decrease in the normalized endocytic flux (Fig. 6b). Note that R1R2(pS-PICK1), which is displayed in Fig. 6c, is not shown in Fig. 6d owing to its lower levels than that of other species. These observations indicate that the competition between the phosphorylation/dephosphorylation of AMPARs at the synaptic plasma membrane affect...
AMPAR trafficking. In addition to the results shown here, we find that the activation levels of PKA and PP2B are not effectively regulated by differences in Ca²⁺ concentration during LTP and LTD stimulation (Supplementary Fig. S7). Thus, although the unbinding of GluA1 from the synaptic membrane is necessary for the LTD induction, the previous model based on only the GluA1-binding to /GluA1-unbinding from the synaptic membrane is insufficient to explain the NMDA receptor-dependent synaptic plasticity. The Ca²⁺-dependent competition between Syt1-mediated exocytosis and PICK1-regulated endocytosis is necessary as the dominant mechanism on LTD as well as LTP.

Dynamics of recycling endosomes during LTP induction and the role of myosin Vb transport. Myosin Vb is widely expressed in most neurons, including those in the hippocampus, which implies the possibility that myosin Vb could mediate endosomal trafficking during LTP induction. Indeed, it has been demonstrated that the genetic chemical inhibition of myosin Vb motility through the binding of nonhydrolysable PE-ADP impairs LTP induction. On the other hand, the basolateral diffusion movement of AMPARs from the dendrites/extrasynaptic area to the synaptic membrane is the dominant mechanism for AMPAR recruitment during LTP. However, in spite of extensively observing numerous experimental evidences, the Ca²⁺-dependent exocytosis pathway of the recycling endosome assisted by myosin Vb transport seems to have lately fallen out of favor as the main pathway for AMPAR trafficking. In fact, recently proposed models for cerebellar LTP and LTD have taken into account the lateral diffusion of AMPARs from the extrasynaptic area to the synaptic membrane as the main pathway of AMPAR trafficking. In addition to the influential arguments that support the long-range lateral diffusion pathway, one of the reasons
for this in the case of hippocampal LTP arises from the following observation: LTP induction is immediately caused by the prompt incorporation of AMPARs into the synaptic membrane, whilst myosin-Vb transport starts several-minutes later than the onset of LTP stimulation, and continues for ~ 10 min. These facts, that seem to contradict each other at first glance, are actually not problematic, at least in the context of hippocampal LTP. In our simulation, the flux from myosin-Vb transport of newly internalized endosomes reaches a maximum at 2 min after the onset of LTP stimulation (Fig. 4a), and continues for a total of ~ 10 min (see the increase in \([R1R2psd(GRIP)\text{MyoVSyt1}]\) in Fig. 5c). Even so, LTP induction can still be reproduced by the network model. Such an apparent discrepancy can be resolved as follows: the prompt incorporation of AMPARs into the synaptic/perisynaptic membranes followed by the rapid diffusional relocation of AMPARs from the peri-synaptic to synaptic membrane, which is required for the immediate induction of hippocampal LTP, is achieved by the Syt1-mediated exocytosis of recycling endosomes localizing on the membrane surface (Fig. 7). This is because the recycling endosomes have already been transported into the synaptic/perisynaptic membrane surface by myosin-Vb in a Ca2+-independent manner. This theoretical prediction is supported by the colocalization of Syt1 with endosomal vesicles and the localization of Syt1 at the peri-synaptic/synaptic membranes. Taking the experimental observations discussed in this study and our simulation results together, we reach a plausible hypothesis that reconciles the long-standing controversy around the AMPAR trafficking pathway: the short-range diffusional relocation pathway of AMPARs exocytosed by Ca2+-dependent Syt1 at the peri-synaptic membranes can primarily contribute to AMPAR recruitment during LTP more than the cell surface long-range lateral diffusion pathway for AMPARs exocytosed at the dendrites/extrasynaptic region. In addition, to reconcile this hypothesis with Penn's observations, we can conclude that the total amount of AMPARs exocytosed at the peri-synaptic membranes should be sufficiently larger than that at the synaptic membrane due to the larger area of peri-synaptic membranes. Therefore, the short-range diffusional relocation pathway of exocytic AMPAR from the peri-synaptic membranes should be more dominant than the direct exocytosis pathway of AMPAR into the synaptic membrane.

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Author contributions
T. S. designed this study, built the computational model, performed the simulations, and analyzed the data. T.S. and K.H. discussed the results. T.S. wrote the manuscript. T.S. and K.H. reviewed and revised the manuscript.

Competing interests
The authors declare no competing interests.

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