Metabotropic Acetylcholine and Glutamate Receptors Mediate PI(4,5)P_2 Depletion and Oscillations in Hippocampal CA1 Pyramidal Neurons in situ

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The sensitivity of many ion channels to phosphatidylinositol-4,5-bisphosphate (PIP_2) levels in the cell membrane suggests that PIP_2 fluctuations are important and general signals modulating neuronal excitability. Yet the PIP_2 dynamics of central neurons in their native environment remained largely unexplored. Here, we examined the behavior of PIP_2 concentrations in response to activation of Gq-coupled neurotransmitter receptors in rat CA1 hippocampal neurons in situ in acute brain slices. Confocal microscopy of the PIP_2-selective molecular sensors tubbyCT-GFP and PLCδ1-PH-GFP showed that pharmacological activation of muscarinic acetylcholine (mAChR) or group I metabotropic glutamate (mGluRI) receptors induces transient depletion of PIP_2 in the soma as well as in the dendritic tree. The observed PIP_2 dynamics were receptor-specific, with mAChR activation inducing stronger PIP_2 depletion than mGluRI, whereas agonists of other Gαq-coupled receptors expressed in CA1 neurons did not induce measurable PIP_2 depletion. Furthermore, the data show for the first time neuronal receptor-induced oscillations of membrane PIP_2 concentrations. Oscillatory behavior indicated that neurons can rapidly restore PIP_2 levels during persistent activation of Gq and PLC. Electrophysiological responses to receptor activation resembled PIP_2 dynamics in terms of time course and receptor specificity. Our findings support a physiological function of PIP_2 in regulating electrical activity.

Phosphatidylinositol-4,5-bisphosphate (PIP_2) directly controls many cellular functions, including membrane and cytoskeletal dynamics and the activity of membrane proteins. These regulatory effects of PIP_2 are mediated by modulation of activity or of membrane association of PIP_2-interacting proteins. In particular, many ion channels are highly sensitive to manipulation of PIP_2 levels.

In cell culture models activation of Gαq-coupled receptors can deplete the PIP_2 content of the plasma membrane by activating phospholipase Cβ (PLCβ) signaling. Such PIP_2 concentration changes were also observed in primary cultures of purkinje and hippocampal neurons. A series of thorough analyses of PIP_2 signaling in response to muscarinic receptor activity in isolated sympathetic ganglion neurons (SGC) established that in these neurons activation of some Gq-coupled receptors leads to transient depletion of PIP_2, which in turn inhibits Kv7.2/3-mediated M-currents. Hence, PIP_2 depletion downstream of receptor-mediated pathways may be a ubiquitous principle controlling neuronal activity by modulating ion channels. Thus it seems likely that the well-known increase of excitability by modulatory neurotransmitters, e.g. in hippocampal pyramidal neurons, is mediated by deactivation of PIP_2-dependent channels. There is also some evidence for PIP_2-dependent regulation of Kir and K2P channels in striatal and thalamic neurons, respectively. However, channel modulation may as well be mediated by other cellular signals downstream of Gq activity. Thus Kv7 channels are also inhibited...
by intracellular Ca\(^{2+}\) elevation\(^{28}\) and inhibition of Gq-sensitive TASK channels is mediated by DAG\(^{27}\). Both, Ca\(^{2+}\) and DAG signals may occur without a substantial drop in PIP\(_2\), downstream of PLC\(^{38-40}\).

Another issue is the method used to assess PIP\(_2\) dynamics. The standard approach is live-cell fluorescence microscopy using genetically encoded sensors built upon PIP\(_2\)-binding domains fused to fluorescent proteins\(^2\). The most popular sensor domain also used by the studies cited above is the pleckstrin homology domain from PLC\(_{δ1}\) (PLC\(_{δ1}\)-PH)\(^{31,32}\). However, interpretation of the observations is complicated by the IP\(_3\) affinity of PLC\(_{δ1}\)-PH, compromising its suitability as a sensor of PIP\(_2\) following PLC activation\(^{30,34}\). More recently, the PIP\(_2\)-specific tubby\(_{CT}\) domain enabled unequivocal measurement of Go\(_q\)-induced PIP\(_2\) depletion in cultured sympathetic and hippocampal neurons\(^{15,33}\). While these findings derived from isolated neurons are consistent with substantial PIP\(_2\) concentration changes, they might still not reflect physiological conditions. Studies in cardiac myocytes showed that PIP\(_2\) content can considerably differ between isolated cells and cells \textit{in situ}\(^{35}\). Besides differences in the extent of PIP\(_2\) depletion, differences in time course may also have relevance for signaling via this pathway. Thus, knowledge of PIP\(_2\) dynamics in native neurons is required for understanding their physiological significance.

To address these issues, we characterized the PIP\(_2\) concentration behavior induced by activation of Go\(_q\) and PLC-coupled transmitter receptors in hippocampal CA1 pyramidal neurons \textit{in situ} in acute brain slices. These neurons receive modulatory cholinergic input from the septohippocampal pathway\(^{36}\), which is mediated postsynaptically by Gq-coupled M1/M3 receptors and results in transient changes of excitability\(^{37-39}\). We find that mAChR and mGluRI receptors induce robust PIP\(_2\) depletion in soma and main apical dendrites. Strikingly, both receptors induced PIP\(_2\) oscillations. Moreover, PIP\(_2\) depletion was receptor and neuron type-specific. Correlation with changes in electrophysiological activity supports an instructive signaling role of these neuronal PIP\(_2\) dynamics.

**Results**

**Muscarinic receptors mediate PIP\(_2\) dynamics in CA1 neurons \textit{in situ}**. We began our investigation of PIP\(_2\) dynamics in acute brain slices by examining the response of hippocampal CA1 pyramidal neurons to activation of their muscarinic ACh receptors. In order to measure Go\(_q\)-induced PIP\(_2\) dynamics \textit{in situ}, we expressed genetically encoded PIP\(_2\) sensors by stereotaxic injection of lentiviral expression vectors into the hippocampi of juvenile (P21) rats (see Methods). Two different GFP-fused sensor domain were used, tubby\(_{CT}\)-GFP\(^{31,32}\) and PLC\(_{δ1}\)-PH-GFP\(^{31,32}\). Both sensors work as ‘translocation sensors’, i.e. their degree of membrane association is a direct measure for PIP\(_2\) concentration and its temporal dynamics\(^3\).

Fluorescence of neurons in acute slices from rats (P26-32) infected with the vector encoding tubby\(_{CT}\)-GFP indicated successful expression in CA1 pyramidal neurons. As shown in Fig. 1a GFP fluorescence was primarily localized to the plasma membrane of the soma and the dendritic tree.

Activation of mAChR receptors by application of the specific agonist, oxotremorine-M (Oxo-M), induced massive and reversible translocation of the tubby\(_{CT}\) probes from the membrane to the cytoplasm in \textit{>95%} of CA1 pyramidal cell somata examined, indicating strong PIP\(_2\) depletion (Fig. 1c). To quantify extent and time-course of probe translocation and hence PIP\(_2\) dynamics we measured fluorescence intensity changes in cytosolic ROIs, as cytosolic signals turned out to be less sensitive to tissue movement than when measuring from the small membrane compartment. Consequently an increase of fluorescence signal corresponds to probe dissociation from the membrane, indicating a loss of PIP\(_2\). Figure 1d shows a representative response to application of Oxo-M for 200 s. The mean tubby\(_{CT}\) translocation reached a peak cytoplasmic amplitude (F/F\(_0\)) of 1.73 ± 0.06 (mean ± SEM; n = 28; 26 slices; 16 rats; Fig. 1e). Mean response latency was 8.8 ± 2.5 s and 90% of the peak response (t\(_{90}\)) was reached within 19.9 ± 2.4 s. Upon washout of the agonist PIP\(_2\) levels recovered within about 100 s as indicated by re-association of the probe to the membrane. Cyttoplasmic fluorescence returned to 10% of peak amplitude (t\(_{10}\), i.e. 90% recovery) in 65.1 ± 7.6 s. To explore the variability of muscarinic PIP\(_2\) dynamics, Oxo-M was applied repetitively with subsequent stimulations separated by a time interval of more than 10 minutes (Fig. 1f). On average, the degree of PIP\(_2\) depletion exhibited a slight but consistent decline in the course of repetitive stimulation. This is consistent with desensitization of muscarinic signaling previously observed in primary hippocampal neurons\(^{11}\).

Previous experiments with cultured neurons have used another sensor domain, PLC\(_{δ1}\)-PH, to examine PIP\(_2\) dynamics\(^9,17,18\). In contrast to the tubby\(_{CT}\) sensor, however, PLC\(_{δ1}\)-PH has a significant affinity for IP\(_3\), which is produced whenever PIP\(_2\) is cleaved by PLC\(_{δ1}\)\(^{33,34,40,42-44}\). Therefore the reliability of PLC\(_{δ1}\)-PH as an indicator of PIP\(_2\) dynamics during GqPCR/PLC signaling has remained an unresolved issue\(^{33,34}\), which provided the rationale for choosing tubby\(_{CT}\) in the present study. Indeed, previous studies showed considerable differences in the behavior of tubby\(_{CT}\)-GFP and PLC\(_{δ1}\)-PH-GFP sensors in terms of translocation following PLC activation\(^{15,34,40}\). Thus, we were interested in comparing both sensors in acute brain slices. When PLC\(_{δ1}\)-PH-GFP was expressed in CA1 neurons (Fig. 1g), stimulation of mAChRs resulted in robust translocation of fluorescence in all neurons examined (Fig. 1h), similar to the results obtained with tubby\(_{CT}\)-GFP (PLC\(_{δ1}\)-PH-GFP: latency 4.9 ± 1.2 s; t\(_{90}\) 22.3 ± 1.9 s; F/F\(_0\) 1.72 ± 0.07; n = 16, 11 slices, 7 rats). However, the recovery was significantly slower compared to tubby\(_{CT}\)-GFP (t\(_{10}\) 132.7 ± 20.8 s; t-test p = 0.008), suggesting that responses of the PH sensor are co-determined by IP\(_3\) production.

**Receptor-specific PIP\(_2\) depletion in CA1 neurons**. Activation of PLC\(_{δ}\) and subsequent hydrolysis of PIP\(_2\) to IP\(_3\) and DAG is the main signaling pathway of Go\(_q\)-coupled receptors. Yet it is not known if PIP\(_2\) depletion is generally associated with the activation of Go\(_q\) coupled receptors other than M1/M3 receptors in central neurons. For example, in sympathetic ganglion neurons activity of muscarinic and purinergic receptors results in a depletion of PIP\(_2\), whereas bradykinin receptors generate IP\(_3\)-dependent Ca\(^{2+}\) signals without substantial changes of the PIP\(_2\) concentration\(^{17,18,29,45,46}\). CA1 pyramidal neurons express various Go\(_q\)-coupled receptors which could potentially induce PIP\(_2\) depletion, including group 1 metabotropic glutamate receptor\(^{47}\), \(Ω_1\)-adrenoreceptor\(^{48}\), bradykinin B\(_2\) receptor\(^49\), Gq-coupled dopamine \(D_1\)-like\(^{-}\) receptor\(^{-}\), histamine H\(_1\) receptor\(^{56,57}\), P2Y\(_2\) receptor\(^{58,59}\), and 5-HT\(_2A/2C\) receptors\(^{60-62}\). We applied specific agonists of group 1 mGluRs, 5-HT\(_2A/2C\) receptors \(Ω_1\) adrenoreceptors, bradykinin...
receptors, Gq-coupled dopamineD1-like receptors, H₁ histamine receptor, or P2Y₁ receptor and monitored PIP₂ concentrations with tubbyCT-GFP. Of these receptors, only mGluRs induced detectable probe translocation indicative for depletion of PIP₂ (Fig. 2a). mGluRI-induced PIP₂ depletion in neuronal somata was consistent across the population of neurons examined (F/F₀ = 1.37 ± 0.05; n = 15; 14 slices from 11 rats; Fig. 2b).

Figure 1. mAChR-mediated PIP₂ depletion in acute brain slices. (a) Representative confocal image of CA1 pyramidal neurons expressing GFP-tagged tubbyCT-GFP probes one week after injection of viral vector. (b) Schematic diagram of hippocampal slice and positioning of the application capillary in relation to pyramidal neurons in the CA1 area. (c) tubbyCT-GFP translocation during mAChR activation. Under resting conditions probes are associated with PIP₂ at the membrane. Application of Oxo-M (10 µM) induced probe translocation to the cytosol indicating PIP₂ depletion, followed by reassociation to the membrane indicating PIP₂ recovery. (d) Representative somatic response to Oxo-M. Shown are cytoplasmatic fluorescence relative to fluorescence before stimulation (F/F₀, upper panel) and a kymograph of probe translocation between membrane and cytosol (lower panel). (e) Average response from 28 individual neurons (26 slices; 16 rats). Application bar is set according to mean response latency. Reduced baseline fluorescence level at the end of the experiments in (d,e) results from photobleaching of GFP. (f) Translocation amplitudes upon repeated Oxo-M application (n = 11, 11, 11, 5, 4, and 3 for successive applications). Responses of individual neurons are shown in grey. (g) Confocal image (maximum intensity projection) of a representative CA1 pyramidal neuron expressing PLCδ1-PH-GFP. (h) Average translocation of PLCδ1-PH-GFP upon Oxo-M application (n = 16 neurons, 11 slices, 7 rats). Scale bars in (a,f): 50 µm. Contrast enhancement of 0.4% and 3% was applied to images shown in (c,d), respectively.
Figure 2. Somatic PIP₂ dynamics are receptor specific. (a) Representative experiment showing PIP₂ depletion in a CA1 soma in response to application of mGluRI agonist DHPG (10 μM) as determined by tubbyCT-GFP translocation. (b) Average PIP₂ concentration changes obtained from 15 cells as in (a) (blue). Response to activation of mAChRs (grey) is redrawn from Fig. 1e for comparison (n = 28). (c) Summary of peak PIP₂ depletion (translocation of tubbyCT-GFP) upon application of specific agonists for the Gq-coupled receptors indicated. Agonists applied were Oxo-M (10 μM, n = 28 neurons/26 slices/16 rats), DHPG (10 μM, n = 15/14/11), DOI, 10–20 μM, n = 14/14/6), methoxamine (10–20 μM; n = 14/14/6), bradykinin (10–20 μM, n = 10/10/4), SKF 83959 (10–20 μM, n = 5/5/3), 2-pyridylethylamin (10–50 μM, n = 10/10/4), ADPβS (10 μM; application for 30 to 60 s, n = 13/13/4). Numbers of experiments also indicated above bars. (d) Representative sensor responses measured from the same CA1 neuron during successive application of 5-HT₂A/₂C agonist DOI and mAChR agonist Oxo-M. (e) Response to higher concentration (100 μM) of DHPG did not increase tubbyCT-GFP translocation (application for 5 min each; n = 6, 6 slices, 3 rats). Successive responses of individual
neurons shown in grey. (f) Average response of PLCδ1-PH-GFP sensor to mGluR1 activation (blue, DHPG) and mAChR activation (black; Oxo-M; replotted from Fig. 1b). (g) Peak translocation of PLCδ1-PH-GFP sensor during application of agonists as in (c); Oxo-M (n = 16 neurons, 11 slices, 7 rats), DHPG (n = 15/11/7), DOI (n = 19/13/7), methoxamine (n = 18/12/7), bradykinin (n = 19/13/7), SKF 83959 (n = 16/11/6), 2-pyridylethylamin (n = 15/11/6), ADPβS (n = 19/13/7). (h) Example responses from two different neurons to application of DHPG (10µM). Note that one response (dark blue) closely matches the average response while the other (light blue) shows pronounced recovery in the presence of the agonist. (i) Responses to repeated DHPG application (n = 11, 11, 3, 3, and 2 for successive applications, respectively; delay between applications ≥ 10 min; individual responses indicated in light blue).

As summarized in Fig. 2c, PIP2 levels in CA1 neurons were insensitive to activation of any of the other receptors. Importantly, subsequent control application of Oxo-M triggered robust translocation of tubbyCT-GFP in each neuron, indicating proper responsiveness of the cell and appropriate sensitivity of the detection approach (Fig. 2d). Further, the prolonged application of each of the agonists (except muscarinic and glutamatergic) for up to 120 seconds or application of the endogenous ligands serotonin and dopamine did not evoke detectable responses (not shown). Equivalent results were obtained with neurons expressing the alternative PIP2 sensor domain, PLCδ1-PH-GFP. As shown in Fig. 2f and g, the stimulation of mAChR and mGluR1 but none of the other receptors examined induced translocation of PLCδ1-PH-GFP. In summary, results obtained with both sensor domains indicate that α1A-adrenoceptor, bradykinin, dopamine2A, histamin-H1, P2Y1, and 5-HT3A/5D do not induce significant PIP2 depletion in the soma of CA1 pyramidal neurons. Thus, PIP2 depletion is specific to mAChR and mGluR1, at least in the context of standard experimental conditions. However, response magnitude of mGluR1 activation was significantly lower compared to muscarinic PIP2 depletion in a population of cells challenged by both agonists (n = 12, 11 slices, 8 rats; paired t-test p = 0.032). While Oxo-M and DHPG have similar binding affinities for their cognate receptors6, EC50 values for downstream effects such as Ca2+ responses are often higher for DHPG than for Oxo-M, raising the possibility that 10µM of DHPG might not be sufficient to evoke a saturating PIP2 response. However, increasing the concentration to 100µM or the duration of agonist application of the glutamatergic agonist (DHPG) did not further increase the response mediated by mGluRs (Fig. 2e) and these responses were significantly smaller than muscarinic responses in the same neurons (p < 0.05, one way ANOVA followed by Tukey post-hoc test, n = 6, 6 slices, 3 rats).

In addition to the smaller responses, mGluR1-induced depletion of PIP2 also differed in its time course compared to muscarinic stimulation (Fig. 2a,b,f). As measured with tubbyCT-GFP, response latencies (6.81 ± 0.89 s) and rise time (t90 = 12.68 ± 1.15 s) were comparable, but time course of recovery was faster compared to mAChR activation (t100 = 28.38 ± 3.57 s; t-test p = 0.0001) Remarkably, in 6 out of 15 recordings, PIP2 levels recovered in the continued presence of the agonist DHPG, as illustrated by individual recordings shown in Fig. 2h. Similarly, PIP2 dynamics as measured with the PLCδ1-PH-GFP probe showed a much faster recovery after activation of mGluR1 (t100 = 30.44 ± 2.05 s, n = 15, 14 slices, 7 rats) when compared to mAChRs (t100 = 132.74 ± 20.77 s; t-test p = 0.0003; Fig. 2f). As noted with mAChR activation, PIP2 dynamics induced by mGluR1s showed slight desensitization in response to repeated application of the agonist (Fig. 2i).

**Dendritic PIP2 dynamics.** Next, we were interested in the spatial pattern of PIP2 depletion, in particular with respect to dendritic compartments. Because we probed PIP2 dynamics with translocation sensors that require microscopic resolution of membrane versus cytoplasm the measurements were confined to dendrites with a diameter of more than 1 μm that were localized close to the slice surface allowing for good optical access. Thus we achieved recordings from the main apical dendrite and its major branches up to 300 μm and basal dendrites to 20 μm distal to the soma. The distance between *stratum pyramidale* to *fissura hippocampi* defining the total length of apical dendrites was about 400 μm for our slices.

We found robust but receptor-specific PIP2 depletion in all dendritic compartments examined. Figure 3a shows a representative example illustrating membrane localization of tubbyCT-GFP in a dendrite and its transient redistribution into the dendritic cytosol during pharmacological activation of mAChRs, indicating reversible depletion of PIP2. With the exception of one apical dendrite (190 μm distance from soma), all dendrites examined (n = 12) responded with the translocation of tubbyCT-GFP. The time course of representative endogenous dendritic PIP2 dynamics is further shown in Fig. 3d as a kymograph and quantitatively as the change of cytosolic fluorescence intensity. The average peak amplitude derived from dendrites of 12 neurons was F/F0 = 1.57 ± 0.06 (12 slices, 10 rats). The average latency of the responses was 14.29 ± 5.26 s. We observed a rise time t90 of 17.52 ± 2.67 s and 90% recovery time (t100) = 41.27 ± 7.25 s after the end of the application. As shown in Fig. 3a,e, activation of mGluR1 also resulted in translocation of tubbyCT-GFP. However, the degree of PIP2 depletion was significantly weaker compared to activation of mAChRs (F/F0 = 1.18 ± 0.02; paired t-test, p = 0.041, n = 3 dendrites in 3 slices from 2 rats). Similar observations were made with PLCδ1-PH-GFP as the PIP2 probe (Fig. 3c,h). Thus, activation of muscarinic receptors by Oxo-M induced strong translocation (F/F0 = 1.86 ± 0.07; t90 = 21.86 ± 1.90 s; t100 = 142 ± 30.53 s; n = 7 dendrites, 7 slices, 3 rats), whereas activation of mGluR1 receptors by DHPG induced small yet reproducible translocation of PLCδ1-PH-GFP (F/F0 = 1.19 ± 0.02; n = 7). Stimulation of various other Gq-coupled receptors did not induce detectable translocation of PLCδ1-PH-GFP (Fig. 3c).

We wondered about spatial, basal-to-apical signaling gradients along the dendrites. Figure 3g,i display amplitudes and time constants of PIP2 depletion as a function of the distance from the soma. We find that neither parameter shows an evident trend along the dendritic length, suggesting that PIP2 depletion is mostly homogenous throughout the larger dendritic compartments examined here (Pearson correlation coefficients:

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**Note:** The text snippet includes scientific data and analysis, describing the responses and dynamics of PIP2 depletion in neurons under various agonist stimulations, with a focus on dendritic compartmental differences. The methods and results are detailed with specific measurements and statistical tests, indicating the receptor specificity and spatial patterns of PIP2 depletion. The data suggest that PIP2 depletion is specific to mAChR and mGluR1 but is not induced by other agonists at the same concentration or duration, highlighting the receptor specificity in cellular signaling responses.
Figure 3. Dendritic PIP$_2$ dynamics. (a) Translocation of tubbyCT-GFP in response to Oxo-M application (10µM) in an apical dendrite 60µm from the soma. (b) Translocation of tubbyCT-GFP from dendritic membranes induced by mAChR (Oxo-M, 10µM, n = 12 neurons/12 slices/10 rats) and mGluRI (DHPG, 10µM, n = 3/3/2) activation, quantified as changes of axial (cytoplasmic) fluorescence changes relative to pre-stimulus fluorescence. (c) Translocation of PLCδ1-PH-GFP from dendritic membranes in response to activation of various Gq-coupled receptors. Agonists used as in Fig. 2 (n = 7 neurons, 7 slices, 3 rats for each condition). (d) Time course of PIP$_2$ depletion from a dendritic recording 220µm from the soma, shown as the relative increase of axial fluorescence (upper panel) and corresponding kymograph (lower panel). (e) Representative fluorescence change and kymograph of dendritic tubbyCT-GFP translocation in response to application of DHPG (10µM; apical dendrite; 20µm from soma). (f) Average time course of mAChR-induced dendritic PIP$_2$ dynamics. (g) Amplitudes (black) and time constants (green) of tubbyCT-GFP translocation in response to application of Oxo-M plotted as a function of the distance of the recording location from the neuronal soma.
Data from 13 individual neurons. (b) Average time course of dendritic PLCδ1-PH-GFP sensor translocation in response to application of the agonists indicated (n = 7 each). (i) Properties of translocation of PLCδ1-PH-GFP in response to Oxo-M plotted as a function of dendritic recording site relative to soma. Contrast enhancement of 10%, 1% and 3% was applied to images shown in a, d and e, respectively.

δlation also revealed considerable variability and complexity in the time course of PIP2 dynamics (Fig. 4b). Most potent PIP2 resynthesis capability of CA1 neurons during receptor activation.

and 16 for mAChR and mGluRI activation, respectively; Fig. 4a). The initial rate of PIP2 decrease was similar for both receptors (t50 = 39.3 ± 5.3 s and 43.3 ± 8.8 s, respectively), but as previously seen with brief receptor stimulation (Fig. 2), the muscarinic responses had a higher average amplitude compared to glutamatergic responses (F/F0 = 1.84 ± 0.08 and 1.55 ± 0.07, respectively; paired t-test p = 0.0002, n = 16, 14 slices; 5 rats). PIP2 recovery after muscarinic receptor activation was more pronounced than for mGluR stimulation such that PIP2 levels tended towards similar values at the end of receptor stimulation period. For both receptors, recovery of PIP2 levels after removal of the receptor agonist was slower than observed with brief receptor stimulation (mACHR: t50 = 243.2 ± 43.9 s; mGluRI: t50 = 167.8 ± 41.5 s; cf. Fig. 2b) and thus depended on the duration of receptor activation.

While demonstrating partial desensitization of PIP2 responses as a common pattern, the prolonged stimulation also revealed considerable variability and complexity in the time course of PIP2 dynamics (Fig. 4b). Most strikingly, PIP2 depletion was often multiphasic or oscillatory. Examples for such complex PIP2 dynamics are shown in Fig. 4c–e. In these cells, PIP2 levels returned to baseline despite sustained presence of the agonist, and moreover, multiple depletion events occurred in rapid succession (Fig. 4d,e). Altogether, 8 out of 17 neurons showed oscillatory PIP2 concentration dynamics with Oxo-M and two out of 16 during application of DHPG. These observations suggest that PIP2 dynamics may be subject to complex temporal regulation and indicate potent PIP2 resynthesis capability of CA1 neurons during receptor activation.

Prolonged receptor activation revealed complexity of PIP2 dynamics. The occasionally observed early recovery of PIP2 level during activation of mGluRI prompted us to examine the time course of PIP2 dynamics during sustained stimulation. Figure 4 shows the resulting PIP2 dynamics measured with the tubbyCT sensor in neuronal somata during continuous application of the receptor agonists for 5 min. Notably, PIP2 depletion generally showed a phasic-ionic time course with an initial strong PIP2 depletion following by partial recovery in the sustained presence of the agonist as apparent from the average from a larger number of neurons (n = 17 and 16 for mACHR and mGluRI activation, respectively; Fig. 4a). The initial rate of PIP2 decrease was similar for both receptors (t50 = 39.3 ± 5.3 s and 43.3 ± 8.8 s, respectively), but as previously seen with brief receptor stimulation (Fig. 2), the muscarinic responses had a higher average amplitude compared to glutamatergic responses (F/F0 = 1.84 ± 0.08 and 1.55 ± 0.07, respectively; paired t-test p = 0.0002, n = 16, 14 slices; 5 rats). PIP2 recovery after muscarinic receptor activation was more pronounced than for mGluR stimulation such that PIP2 levels tended towards similar values at the end of receptor stimulation period. For both receptors, recovery of PIP2 levels after removal of the receptor agonist was slower than observed with brief receptor stimulation (mACHR: t50 = 243.2 ± 43.9 s; mGluRI: t50 = 167.8 ± 41.5 s; cf. Fig. 2b) and thus depended on the duration of receptor activation.

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Modulation of electrical behavior by receptor stimulation. The well-described effects of muscarinic activity on the electrical properties of CA1 neurons – including inhibition of M-currents – may (at least partially) be mediated by PIP2 concentration dynamics. We thus were interested in differential effects on excitability of the various Gq/PLC-coupled receptors examined for their coupling to PIP2 dynamics.

To this end, we performed patch clamp experiments in current clamp mode in acute brain slices prepared from rats at P14 to P21. Current step protocols were used to assess membrane potential, input resistance, spiking behavior, and afterpolarisation (Fig. 5a). Experiments were performed in the presence of inhibitors of GABA_A/B and ionotropic glutamate receptors (see Methods) to exclude effects resulting from network activity.

Overall, we found pronounced changes in electrical behavior following the activation of muscarinic mACHR receptors and mGluRI but little effects of other Gq-coupled receptors. Consistent with previous findings agonists of both mACHR (n = 11, 11 slices, 10 rats) and mGluRI (DHPG 10 µM, n = 9, 8 slices, 5 rats) induced depolarization of the resting membrane potential and an increase in firing frequency during depolarization (number of action potentials, NAP; Fig. 5a–d,f). In CA1 cells, a train of action potentials is usually followed by an afterhyperpolarization (AHP) 37,68,69. Application of either Oxo-M or DHPG resulted in the disappearance of the AHP and the appearance of an afterdepolarisation (ADP; Fig. 5a,e). All of these receptor-induced changes are consistent with the deactivation of potassium conductances such as M currents 37,68,70,71. In some neurons, activation of mACHR receptors induced sustained depolarization (plateau potentials) subsequent to the 600 ms current step, as also described previously 72.

As shown in Fig. 5c–f, stimulation of other Gq-coupled receptors, including 5-HT2A/C receptors, α1 adrenergic receptors, bradykinin receptors, D1-like dopamine receptors, H1 histamine receptors, and purinergic P2Y1 receptors had no significant effects on any of these electrophysiological characteristics. Noteworthy, the mACHR and mGluRI induced depolarization did not always persist during the agonist application, as occasionally transient hyperpolarisation and oscillations of the membrane potential was observed during mACHR activation. During muscarinic stimulation, 9 of 11 neurons showed substantial recovery from depolarization in the presence of Oxo-M; of those, 6 neurons showed complete repolarization or even hyperpolarisation before washout. In the presence of DHPG, 3 neurons showed a substantial recovery from initial depolarization. Oscillatory behavior was observed in neurons from younger animals (P14-21), but also in slices age-matched to the PIP2 imaging experiments, as shown in an exemplary recording (Fig. 5g) obtained from a neuron in a P27 slice.

In summary, we find that pronounced changes in membrane potential and firing rates paralleled neuronal PIP2 depletion in terms of effect size, time course and receptor specificity. Thus, depolarization, increased spike rates and PIP2 dynamics were largely restricted to the activation of mACHR and mGluRI receptors.
Neuron type-specific PIP₂ dynamics: dentate gyrus granule cells. To extend our observations on PIP₂ dynamics during extended application of mAChR and mGluRI agonists Oxo-M (n = 17 neurons/15 slices/5 rats) and DHPG (n = 16/14/5) as measured by translocation of tubbyCRT-GFP. (b) Distribution of distinct temporal patterns of tubbyCRT-GFP sensor translocation within the population of neurons challenged by sustained application (5 min) of agonists. Responses were categorized as constant, desensitizing (partial recovery), transient (full recovery) and oscillating (≥2 peaks). (c) Distinct temporal behavior of PIP₂ dynamics of an individual pyramidal neuron to successive applications of Oxo-M and DHPG. Lower panel shows a kymograph illustrating the response to Oxo-M. (d) Oscillatory PIP₂ dynamics in a CA1 neuron in response to prolonged activation of mACh receptors. Note the pronounced repetitive phases of rapid PIP₂ depletion and replenishment during stimulation. (e) Oscillatory PIP₂ dynamics of a neuron in response to continuous activation of mGluRs. Contrast enhancement of 0.4%, 1% and 5% was applied to kymographic images shown in c, d and e, respectively.

Discussion

Direct observation of PIP₂ dynamics in central neurons in situ. While there is good evidence for PIP₂ depletion in response to activation of Gq-coupled receptors for some types of neurons studied in the cell culture dish, surprisingly little is known about the prevalence and spatiotemporal properties of PIP₂ dynamics in central neurons under physiological conditions. PIP₂ levels and their dynamic regulation may be largely different in vivo, as embedding in the native environment and full differentiation of neurons may impact on relevant factors such as expression and spatial subcellular organization of receptors and downstream components of the signaling cascade and the enzymes that resynthesize PIP₂. Therefore, information on PIP₂ concentration behavior in intact tissue preparations such as brain slices is required. Previous studies with organotypic slices were consistent with PIP₂ depletion in situ triggered by synaptic release of glutamate onto cerebellar Purkinje neurons or by muscarinic agonist application in cortical pyramidal cells. However, both studies were not fully conclusive with
Figure 5. Electrophysiological responses to activation of Gq-coupled receptors. (a,b) Current clamp protocol (lower panel) and representative recordings of corresponding membrane potential (V_m) responses of CA1 pyramidal neurons before (light traces) and during (dark traces) activation of mAChRs or mGluRIs by 10µM Oxo-M or DHPG, respectively. (c) Changes of resting membrane potential (ΔV_m) displayed as the difference before and upon application of various GqPCR activators. Agonists applied were Oxo-M (10µM; n = 10 neurons/10 slices from 7 rats; data points with plateau potentials were excluded for this analysis) for AChRs, DHPG (10µM; n = 9/8/5) for mGluRIs, DOI (20µM; n = 8/8/7) for 5-HT2A/C, methoxamine (20µM; n = 8/8/7) for α1-adrenergic receptors, bradykinin (20µM; n = 7/7/6), SKF 83959 (20µM; n = 7/7/6) for D1-like dopamine receptors, 2-pyridylethylamin (20µM; n = 8/8/7) for H1 histamine receptors, ADPβS (10µM; n = 15/15/11) for P2Y1-R. Asterisks indicate significance of difference to control application of ACSF (n = 10/10/7) with p < 0.05 (*), 0.01 (**) and 0.001 (***) (one-way ANOVA followed by Dunnett multiple comparison test). (d) Difference in number of action potentials (AP) triggered during 600 ms depolarizing current step. Numbers of experiments as in (c). (e) Changes in afterpolarisation where a negative Δ indicates an AHPm increase and positive Δ indicates an AHP reduction or afterdepolarisation. Numbers of experiments as in (c). (f) Mean time courses of V_m modulation during application of ACSF as a control (grey), Oxo-M (black), DHPG (blue), and DOI (purple). V_m was measured at the first 500 ms of each trace in the absence of spiking; see (a). (g) Example membrane voltage oscillation during mAChR activation in a P27 slice.
respect to PIP$_2$ signaling because PLC$\delta$-PH was used as a sensor domain, which has a similar affinity for PIP$_2$ and IP$_3$ and may report the production of IP$_3$ rather than depletion of PIP$_2$. In fact, Okubo et al. interpreted probe translocation in terms of IP$_3$ production rather than depletion of PIP$_2$. Here, by using tubby$\textsubscript{CT}$, as an alternative PIP$_2$ sensor insensitive to IP$_3$, our present data now show unequivocally that muscarinic and metabotropic glutamate receptors indeed trigger PIP$_2$ dynamics in a prototypic central neuron in situ. Of note, in cultured cell lines tubby$\textsubscript{CT}$ previously failed to respond or only weakly translocated upon PLC-mediated PIP$_2$ depletion, which initially was attributed to a higher affinity for PIP$_2$ compared to PLC$\delta$-PH. However, quantitative titration of PIP$_2$ in living cells showed that its affinity to PIP$_2$ is actually lower, which should make it a useful PIP$_2$ sensor. In fact, our current results demonstrate that tubby$\textsubscript{CT}$ readily translocates in a neuronal cellular environment. The difference in behavior between experimental conditions is not yet understood, but may suggest cell type-specific segregation of PIP$_2$ into distinct pools selectively accessible to the different PIP$_2$-binding domains. In any case, our findings show that in the native neuronal system tubby$\textsubscript{CT}$ is a much better reporter of PIP$_2$ dynamics than might have been anticipated. Thus, using tubby$\textsubscript{CT}$-GFP allowed us to systematically assay PIP$_2$ dynamics without confounding effects of IP$_3$ dynamics.

**Spatiotemporal properties of PIP$_2$ dynamics.** We found that in the larger compartments accessible to measuring sensor translocation, receptor-induced PIP$_2$ depletion appeared largely homogenous without evidence for substantial subcellular differences. This suggested that induced PLC activity is similar in somatic and dendritic compartments. This finding was not entirely expected, because while M1 receptors show high densities throughout soma and dendrites, the most prominent mGluRI receptor of CA1 neurons, mGluR$_5$, has a relatively low density in soma compared to dendrites. Since glutamatergic PIP$_2$ depletion in dendrites was not stronger than in the somatic compartment, the PIP$_2$ depletion pattern does not appear to correlate closely with receptor distribution. It is worth noting that the moderate degree of PIP$_2$ sensor translocation in dendrites did not result from the distinct dendritic geometry, because the smaller volume-to-membrane area ratio in the dendrites should rather result in a stronger relative increase of sensor fluorescence when sensors dissociate from the membrane. Also, muscarinic stimulation elicited larger dendritic responses than glutamatergic stimulation, showing that PIP$_2$ sensor response was not saturated by mGluR stimulation. Of note, a similar observation was made by Nakamura et al. for Ca$^{2+}$ dynamics in CA1 pyramidal cells: activation of mGluRI, mAChR and 5-HT$_2$R elicited comparable Ca$^{2+}$ waves despite different receptor distribution. Thus, these neurons may possess mechanisms to globalize Gq signaling including PIP$_2$ depletion.

To our knowledge our results for the first time demonstrate oscillations of the PIP$_2$ concentration in a neuron. While IP$_3$, DAG and Ca$^{2+}$ are known to undergo oscillatory concentration dynamics in neurons, previous observations on PIP$_2$ dynamics in primary dissociated neurons seemed to indicate that PIP$_2$ concentrations essentially remained depleted during prolonged receptor activity. Oscillatory translocation of the PLC$\delta$-PH sensor domain observed occasionally has been understood as dynamics of the IP$_3$ signal picked up by the PH
δ1-PH and tubby CT as a major cellular signal in the control of neuronal activity through regulation of PIP2-sensitive ion channels such as Kv7. Inhibition of Gq signaling by, e.g., PKC, receptor kinases, or RGS molecules may contribute to a negative feedback loop controlling PIP2 degradation. Moreover, our observations reveal an impressive capability of PIP2 replenishment, as indicated by the rapid and complete recovery of PIP2 levels in presence of agonists. PIP2 resynthesis may be increased during GqPCR activation, providing negative feedback to PIP2 depletion and possibly contributing to observed oscillations. Specifically, PIP2 replenishment may involve Ca2+ and phosphatidic acid-dependent phospholipid exchange at plasma membrane-endoplasmic reticulum (PM-ER) junctions.

Whatever the mechanism underlying the oscillations is, our findings indicate that PIP2 dynamics may provide neurons with another dimension of effector modulation beyond a simple on/off switch for downstream effectors. Although the consequences of PIP2 oscillations for electrical neuronal activity remain to be explored, we note that indeed, neurons showed fluctuations of membrane potential and firing frequency during agonist application. It is worth mentioning that mACHR and mGluRI agonists can induce and shift gamma and theta oscillations. In the light of the present data, it is tempting to speculate that PIP2 oscillations might participate in such frequency modulation.

Neuronal ion channels as effectors of PIP2 dynamics. Given the known high sensitivity of some ion channels to even a moderate drop in the PIP2 concentration, a main potential target of PIP2 depletion are ion channels and thus electric excitability. Based on studies on isolated neurons, inhibition of Kv7 channels as the direct consequence of PIP2 depletion is well established. Our data permit the correlation of PIP2 dynamics and electrophysiology in situ. Activation of mACHR and mGluRI, but not other PLC-coupled receptors known to be present and functional in CA1 neurons induced robust PIP2 depletion. The same pattern of receptor specificity was observed for modulation membrane potential, firing frequency and afterhyperpolarization, providing at least circumstantial evidence for the causation of channel regulation by PIP2. Simultaneous recordings of electrical activity and PIP2 dynamics from the same neuron should be performed in the future to provide more direct evidence.

In conclusion, our data support and generalize the as yet largely hypothetical mechanism of PIP2 dynamics as a major cellular signal in the control of neuronal activity through regulation of PIP2-sensitive ion channels such as Kv7. Future studies need to address this issue rigorously by manipulating PIP2 levels in situ. Along those lines a recent study aimed at PIP2 depletion in hippocampal slice cultures by chemically induced recruitment of a PIP2 phosphatase. While this approach did not reveal any effects on electrical properties of the neurons, the results appear inconclusive since changes in PIP2 concentration were not verified.

One of the most intriguing unknowns are the spatiotemporal properties of PIP2 dynamics during entirely physiological neuronal activity, i.e., during synaptic activity of the modulatory (e.g., cholinergic) and principal (i.e., glutamatergic) inputs into the hippocampal neurons and of the PIP2 dynamics associated with intrinsic neuronal (network) activity. Another question is the PIP2 oscillations in the distal smaller dendritic compartments not amenable to analysis by the translocation probes used in this study. In particular, in the immediate postsynaptic compartment, i.e. spines, PIP2 may have a role in controlling synaptic plasticity.

Materials and Methods

Virus production and constructs. Lentiviral plasmids pCMVΔR8.9, pVSVG and FUGW were kindly provided by Pavel Osten (MPI for medical research Heidelberg, Germany). The PLCΔ1-PH and tubbyCT constructs were provided by Tamás Balla (NIH, Bethesda, USA) and Lawrence Shapiro (Columbia University, USA), respectively. Lentiviral particles were derived by triple transfection of HEK293FT cells with Lipofectamin 2000 (Invitrogen, Darmstadt, Germany). Virus purification from supernatant was achieved by 15 minute centrifugation at 3000 rpm, filtration through a Millex® HV 0.45 μm filter (Millipore, Darmstadt, Germany) and two successive ultracentrifugation steps (25000 rpm, 1 h 30 min, 4°C). Pellets were resuspended in TBS-5 buffer (50 mM Tris-HCl, 130 mM NaCl, 10 mM KCl, 5 mM KCl) and subjected to a final 30 centrifugation at 5000 rpm. Aliquots were stored at −80°C and thawed up to two times.

Animals, stereotactic injection and slice preparation. Wistar rats were obtained from the animal facility of the Philippus University of Marburg (Marburg, Germany) or Charles River (Cologne, Germany) and kept and handled according to German law and institutional guidelines at the Philippus University. All procedures were approved by the Regierungspäsidium Giessen, Germany. Animals were housed with access to ad libitum water and food on a 12-h light/dark cycle. At weaning (postnatal day 21) male and female rats were anesthetized with Isoflurane (Baxter, Unterschleißheim, Germany) or Sevorane (Sevorane®, Abbott,
both to the bath and local perfusion. 7 seconds. In applications of the P2Y1 agonist ADP action potential threshold to quantify spiking behavior and afterpolarisation (see Fig. 5a,b). Sweep length was stored in voltage clamp mode before and after each current clamp recording, but not corrected for. Measurements bleaching generally being more pronounced for tubby CT-GFP than for PLC

Medium afterhyperpolarisation (AHP m) was obtained as the difference of resting Vm and mean Vm at 70 to 120 ms Na2-ATP 2, Na2-GTP 0.3, HEPES 10 and EGTA 0.1 (adjusted to pH 7.2 with KOH). Series resistance was monitored in voltage clamp mode before and after each current clamp recording, but not corrected for bleaching which results in apparently lower signals following transient depletion of PIP 2, with fluorescence signal and normalized to signal at the onset of agonist application. We found that probe translocation without evident response were corrected for photobleaching according to a biexponential fit to the decaying fluorescence signal and normalized to signal at the onset of agonist application. We found that probe translocation generally presented the reliable determination of the time course of photobleaching. Therefore most data were not corrected for bleaching which results in apparently lower signals following transient depletion of PIP2, with bleaching generally being more pronounced for tubbyCT-GFP than for PLC1-PH-GFP. Confocal images were further analyzed with ImageJ (National Institutes of Health, USA) to isolate individual images of a time series, create kymographs and set scale bars. Electrophysiological data were recorded with a HEKA EPC10USB amplifier and Patch Master software (Version 2.43 HEKA, Lambrecht, Germany) in current clamp mode. Data were low pass filtered with a 2.9 kHz Bessel filter and digitized at 20 kHz. Borosilicate recording pipettes had a resistance of 3–4 MΩ and were filled with intracellular solution containing (in mM): K-gluconate 135, KCl 20, MgCl 2, Na2-ATP 2, Na2-GTP 0.3, HEPES 10 and EGTA 0.1 (adjusted to pH 7.2 with KOH). Series resistance was monitored in voltage clamp mode before and after each current clamp recording, but not corrected for. Measurements with a change in series resistance >40% during the course of the experiment were discarded. Input resistance was assessed by injection of small positive and negative currents steps, followed by a depolarizing current step above action potential threshold to quantify spiking behavior and afterpolarisation (see Fig. 5a,b). Sweep length was 7 seconds. In applications of the P2Y1 agonist ADP/PS a shorter protocol without the positive 20 pA step was used. Medium afterhyperpolarisation (AHP m) was obtained as the difference of resting Vm and mean Vm at 70 to 120 ms after the depolarizing current step. Changes in AHP value resulting from application of receptor agonists are given as Δafterpolarization such that positive values indicate reduction of AHP or eventually the emergence of an afterdepolarisation. Amplitudes were calculated from averaging at least 10 baseline data points and a minimum of 3 peak points, with avoidance of plateau potentials.

Statistical analysis. Statistical significance was tested in Igor Pro. Randomness, equal variances and normal distribution of the data was tested with Igor’s Runs, Kolmogorov-Smirnov and Jarque-Bera test. In cases where validity of a parametric test was compromised, a Wilcoxon-Mann-Whitney test was performed. Where applicable, groups of two were compared with paired and unpaired Student’s t. Two tailed one-way ANOVA was followed by a Dunnett test for comparing multiple groups to a single control or a Tukey test to compare all groups to each other. Unless noted otherwise all values are given ± standard error of the mean.

Chemicals and perfusion system. OXOteremine-M, DHPG, Bradykinin, SKF83959, DOI, Serotonin and Dopamine were purchased from Tocris and Methoxamine and 2-Pyridylethylamine from Sigma. All other chemicals were from Sigma/Fluka or Merck (Germany). For application of test substances a capillary of 200 μm transversal slices were cut with a vibratome (VT1200, Leica Biosystems, Wetzlar, Germany) and placed into a chamber with 4°C sucrose-ACSF. After a 35 min recovery period at 35°C slices were kept at room temperature. For recordings slices were transferred to a submerged chamber and perfused with ACSF (in mM: 125 NaCl, 25 NaHCO3, 25 D-glucose, 2.5 KCl, 2 CaCl2 and 1 MgCl2, oxygenated with 95% O2/5% CO2) for at least 20 minutes.

Imaging, electrophysiological recording and data analysis. Confocal imaging was performed with a Zeiss LSM710 (Zeiss, Oberkochen, Germany). The sampling rate for time series experiments was 1.75 s with a pixel size of 0.13 μm. In some cases (especially dendrite measurements) the sampling rate was increased to 1 s. In all cases where the sampling rate slightly deviated the data were resampled to allow averaging across experiments. Overlay with the original was performed to ensure preservation of time scale. Average cytoplasmic fluorescence intensities were determined from regions of interest (ROI) excluding both the plasma membrane (defined as the local intensity max at the cell’s border in the resting cell) and the nucleus. Distance of ROIs to the plasma membrane was >0.5 μm even when slight shifts of the cell’s position occurred during the experiment. ROIs were defined post-hoc using the microscope software ZEN (Versions 2008 and 2009) and obtained average intensities were exported to Igor Pro (Version 6.03 A, Wave Metrics, Portland, OR USA). Traces were background subtracted and normalized to the last time point before beginning of a response (F/F0 normalized to t0). Measurements without evident response were corrected for photobleaching according to a biexponential fit to the decaying fluorescence signal and normalized to signal at the onset of agonist application. We found that probe translocation generally presented the reliable determination of the time course of photobleaching. Therefore most data were not corrected for bleaching which results in apparently lower signals following transient depletion of PIP2, with bleaching generally being more pronounced for tubbyCT-GFP than for PLC1-PH-GFP. Confocal images were further analyzed with ImageJ (National Institutes of Health, USA) to isolate individual images of a time series, create kymographs and set scale bars. Electrophysiological data were recorded with a HEKA EPC10USB amplifier and Patch Master software (Version 2.43 HEKA, Lambrecht, Germany) in current clamp mode. Data were low pass filtered with a 2.9 kHz Bessel filter and digitized at 20 kHz. Borosilicate recording pipettes had a resistance of 3–4 MΩ and were filled with intracellular solution containing (in mM): K-gluconate 135, KCl 20, MgCl 2, Na2-ATP 2, Na2-GTP 0.3, HEPES 10 and EGTA 0.1 (adjusted to pH 7.2 with KOH). Series resistance was monitored in voltage clamp mode before and after each current clamp recording, but not corrected for. Measurements with a change in series resistance >40% during the course of the experiment were discarded. Input resistance was assessed by injection of small positive and negative currents steps, followed by a depolarizing current step above action potential threshold to quantify spiking behavior and afterpolarisation (see Fig. 5a,b). Sweep length was 7 seconds. In applications of the P2Y1 agonist ADP/PS a shorter protocol without the positive 20 pA step was used. Medium afterhyperpolarisation (AHP m) was obtained as the difference of resting Vm and mean Vm at 70 to 120 ms after the depolarizing current step. Changes in AHP value resulting from application of receptor agonists are given as Δafterpolarization such that positive values indicate reduction of AHP or eventually the emergence of an afterdepolarisation. Amplitudes were calculated from averaging at least 10 baseline data points and a minimum of 3 peak points, with avoidance of plateau potentials.

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Data availability. Most data generated or analysed during this study are included in this published article. Additional datasets generated and analysed during the current study are available from the corresponding authors on reasonable request.

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**Author Contributions**

D.O. and S.H. designed the study. S.H. performed the experiments and analyzed the data. S.H. and D.O. wrote the manuscript. All authors approved of the manuscript.

**Additional Information**

**Competing Interests:** The authors declare no competing interests.

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