ABSTRACT: Synaptic cooperation and competition are important components of synaptic plasticity that tune synapses for the formation of associative long-term plasticity, a cellular correlate of associative long-term memory. We have recently reported that coincidental activation of weak synapses within the vicinity of potentiated synapses will alter the cooperative state of synapses to a competitive state thus leading to the slow decay of long-term plasticity, but the molecular mechanism underlying this is still unknown. Here, using acute hippocampal slices of rats, we have examined how increasing extracellular dopamine concentrations interact and/or affect electrically induced long-term potentiation (LTP) in the neighboring synapses. We demonstrate that D1/D5-receptor-mediated potentiation at the CA1 Schaffer collateral synapses differentially regulates synaptic co-operation and competition. Further investigating the molecular players involved, we reveal an important role for extracellular signal-regulated kinases-1 and 2 (ERK1/2) as signal integrators and dose-sensors. Interestingly, a sustained activation of ERK1/2 pathway seems to be involved in the differential regulation of synaptic associativity. The concentration-dependent effects of the modulatory transmitter, as demonstrated for dopaminergic signaling in the present study, might offer additional computational power by fine tuning synaptic associativity processes for establishing long-term associative memory in neural networks. © 2015 The Authors Hippocampus Published by Wiley Periodicals, Inc.
Hippocampal DA acting through D1Rs is proposed to mediate informational saliency and thus promote the persistence of long-term memories (Hansen and Manahan-Vaughan, 2014). The hippocampus and VTA connections form a loop that modulates encoding and integration of motivationally relevant, novel, and reward-related information (Lisman and Grace, 2005). Additionally, hippocampal D1R activation modulates the locomotor activity and also dopamine efflux in the nucleus accumbens (Zornoza et al., 2005), thus extending its significance to addiction and reward-related mechanisms.

Altered dopaminergic signaling and modulation has been implicated in the pathophysiology of many neuropsychiatric disorders and also in the age-associated memory deficits. Consequently, administration of dopamine agonists has been reported in many cases to alleviate or restore the alterations. However, dopaminergic signaling is known to exhibit dose-dependent effects in many brain regions with linear or nonlinear profiles (Williams and Millar, 1990; Zheng et al., 1999; Chao et al., 2002). In prefrontal cortex, low levels of D1R activation is proposed to serve a role in sculpting neural selectivity by increasing neural signal-to-noise ratio (Puig and Miller, 2012). However, whether the functional response of dopaminergic signaling differs over a wide range of DA concentrations in hippocampus remains to be explored. Interestingly, bath application of D1R agonists or DA itself induces a slow-onset potentiation in acute hippocampal slices in vitro that occludes electrically induced LTP (Huang and Kandel, 1995; Navakkode et al., 2012). This raises important concerns on how the changes in the extracellular dopamine concentrations influence the ongoing information processing and affect the encoding and maintenance of memories. Here, we have investigated whether different degrees of D1R activation could lead to differential outcomes in the context of late-associative mechanisms. Given that D1R-activation leads to robust activation of mitogen-activated protein kinases (MAPKs) in CA1 region (Roberson et al., 1999), we have also explored their potential role in this context.

### Preparation of Hippocampal Slices

A total of 181 acute hippocampal slices prepared from 125 male Wistar rats (6–7 week old) were used in the study. All animal procedures were approved by the Institutional Animal Care and Use Committee (IACUC) of National University of Singapore. Briefly, after anaesthetization using CO₂, the rats were decapitated and the brains were quickly removed to the cold (2–4°C) aCSF being continuously bubbled with carbogen. Transverse hippocampal slices of 400 μm thickness were prepared from the right hippocampus using a manual tissue chopper (Stoelting, Wood Dale, Illinois), transferred onto a nylon net in an interface chamber (Scientific Systems Design, Ontario, Canada) and incubated at 32°C at an aCSF flow rate of 1 mL/min and carbogen consumption of 16 l/h. The whole process, from anaesthetization to the transfer of slices to the chamber, was carried out very quickly with an average duration of 5 minutes. The slices were incubated for at least 3 hours before starting the experiments.

### Field Potential Recordings

Monopolar, lacquer-coated, stainless steel electrodes (5 MΩ: AM Systems, Sequim, Washington) were used as stimulating and recording electrodes. Stimulating electrodes were positioned within the stratum radiatum of the CA1 region to stimulate Schaffer collaterals and the field-EPSP responses were recorded with an electrode placed in the CA1 distal apical dendritic region. After the preincubation period of at least 3 h, an input–output curve (stimulus intensity vs. fEPSP slope) was plotted for each input and the test stimulus intensity was set to obtain a fEPSP slope 40% of the maximal response. The signals were amplified by a differential amplifier, digitized using a CED 1401 analog-to-digital converter (Cambridge Electronic Design, Cambridge, UK) and monitored online with custom-made software.

In all experiments, a stable baseline was recorded for at least 30 minutes before chemical or electrical LTP induction. Four 0.2-Hz biphasic, constant current pulses (spaced at 5s) given every five minutes were used for baseline and post-induction recording and the average slope value from the four sweeps was considered as one repeat while used for plotting. For two- and three-input synaptic tagging and capture experiments, the stimulating electrodes were placed on either side of the recording electrode so as to stimulate two or three independent set of synapses respectively. A paired-pulse facilitation protocol (Li et al., 2014; Sajikumar et al., 2014) was used to test the independence of the two inputs. Initial slopes of fEPSPs were expressed as percentages of baseline averages.

### Statistics

The time-matched, normalized data were averaged across replicate experiments and expressed as mean ± SEM. The average percentage values of fEPSP slope per time point were subjected to statistical analysis with GraphPad Prism 6.0. Whenever the data did not conform to Gaussian distribution, nonparametric tests were used. Wilcoxon matched-pairs signed rank test was used when comparisons were made within group. Multiple, between group comparisons for specified time-points were performed with either one-way or two-way ANOVA with Tukey’s or Dunnet’s post-hoc tests. Statistical significance was assumed at P < 0.05. (*) P < 0.05 ** P < 0.01 *** P < 0.001, **** P < 0.0001.
LTP Induction Protocols

For early-LTP induction, a weak tetanization protocol (WTET) consisting of a single high frequency stimulation (100 Hz, 21 biphasic constant current pulses, single burst, 0.2 ms pulse duration) was used. For late-LTP induction, a strong tetanization protocol (STET) involving repeated high-frequency stimulation (three trains of 100 Hz, 100 pulses, single burst, 0.2 ms pulse duration) was used with an intertrain interval of ten minutes.

Drugs

The D1/D5-receptor agonist SKF-38393 hydrochloride [(±)-1-Phenyl-2,3,4,5-tetrahydro-(1H)-3-benzazepine-7,8-diol hydrochloride] (SKF; #D047, Sigma-Aldrich, Singapore) was stored at −20°C as 15–50 mM stock in DMSO. Another agonist 6-Bromo-APB hydrobromide [R(+)–6-Bromo-7,8-dihydroxy-3-allyl-1-phenyl-2,3,4,5-tetrahydro-1H-3-benzazepine hydrobromide] (APB; #B135, Sigma-Aldrich, Singapore) was stored as 15–50 mM stock in DMSO at −20°C. The stocks were stored for not more than a week. Just before application, the stocks were diluted to the final concentration in aCSF, bubbled with carbogen, and bath applied for three 5-minute durations with 5-minute interval between each application. The drugs were protected from light during storage and bath application was carried out under dark conditions.

The MEK inhibitor U0126 (#V1121, Promega, Madison, Wisconsin) was stored as 20 mM stock in DMSO just before application. Both the MEK inhibitors were applied after one hour from the start of SKF application and kept throughout the rest of recording period.

Whenever the drug stock was prepared in DMSO, the DMSO concentration in the final bath-application solution was kept below 0.1%, a concentration which has been shown to not affect basal synaptic responses (Navakkode et al., 2005).

RESULTS

Different Concentrations of D1/D5-Receptor Agonists Induce Slow-Onset Potentiation in CA1 Pyramidal Neurons

Dopamine or D1R agonists have been shown to induce a slow-onset potentiation of field-EPSP slopes in hippocampal CA1 region (Gribbolk and Ashe, 1984; Frey et al., 1993; Huang and Kandel, 1995). First series of experiments involved the application of a selective D1R agonist SKF-38393 (SKF) at a range of concentrations from 50 µM to 0.1 µM. We chose these concentrations as many of the studies have generally used a concentration of 50–100 µM (Huang and Kandel, 1995; Roberson et al., 1999; Navakkode et al., 2007; Navakkode et al., 2012) and we were interested to know whether lower concentrations are also capable of inducing a persistent potentiation. As a control experiment, first we recorded stable baseline potentials by delivering low-frequency test stimulation every five minutes in synaptic input S1 (Fig. 1b). The baseline response remained stable over 3 hours. Though a small drift in the responses was observed between 120 and 140 minutes, it was not statistically significant at any time point (P > 0.05, n = 6). In the next series of experiments, after a stable baseline recording of at least 30 minutes, SKF was applied in three 5-minute durations with 5-minute interval between applications. This application paradigm is analogous to the spaced strong tetanization protocols used to induce late-LTP (Navakkode et al., 2007).

Application of 50 µM SKF resulted in a slow-onset potentiation similar to earlier reports (Huang and Kandel, 1995; Navakkode et al., 2007). The potentiation became significantly different from the baseline 35 minutes after the first application (118.1 ± 3.06%, P < 0.05) and maintained over 4 hours. The mean potentiation at the end of 4 hours was (155.12 ± 7.28%) (n = 6, Fig. 1c). With 25 µM SKF application also a slow-onset potentiation was observed that became significantly different from baseline by 25 minutes [(116 ± 2.84%) P < 0.01]. The mean potentiation at the end of 4 hours was (148.77 ± 5.52%) (P < 0.01, n = 10, Fig. 1d). Similar was the result with 10 µM SKF application; the potentiation was significantly different after 20 minutes (114.2 ± 5.78% P < 0.01) and stayed significant for 4 hours (132.03 ± 4.71%, P < 0.01; n = 8, Fig. 1e). Interestingly, even with 5 µM SKF application, a late-onset potentiation was observed that became significantly different rather slowly by 90 minutes (115.05 ± 5.63%, P < 0.05) and stayed significant thereafter till the end of recording [(126.30 ± 8.20%) P < 0.05, n = 8, Fig. 1f]. Application of further lower concentrations of SKF, 1 µM and 0.1 µM, failed to induce any significant potentiation; the mean potentiation at the end of 4 hours being (107.52 ± 3.78%) (P > 0.05, n = 8, Fig. 1g) and (98.71 ± 8.23%) (P > 0.05, n = 8, Fig. 1h) respectively.

A plot of the mean potentiation induced by the different concentrations of SKF at the end of 4 hours revealed a concentration-dependent response (Fig. 1i). A one-way analysis of variance (ANOVA) comparing these normalized group means to baseline (100%) showed a significant difference between the mean values (F(6, 51) = 16.02; P < 0.0001). Multiple comparisons with the Dunnett’s post-test showed that potentiation was significantly different from the baseline with 5 µM (P < 0.01), 10 µM (P < 0.001), 25 µM (P < 0.0001) and 50 µM (P < 0.0001) groups whereas not with 1 µM (P > 0.05) and 0.1 µM (P > 0.05) groups.

We repeated these experiments with another structurally different D1R analogue, 6-bromo-APB (APB), and obtained similar concentration-dependent response (Supporting Information Fig. 1a–g).

Taken together, these series of experiments showed that the D1R-mediated potentiation at the Schaffer collateral-CA1 synapses is concentration-dependent.
FIGURE 1.
Dose-Dependent Regulation of Synaptic Cooperation and Competition by Dopaminergic Signaling

Having observed the concentration-dependent response of the D1R-mediated potentiation, next we were interested to know whether such effects might have any significance during the associative information processing at the input level.

To investigate this, we designed a series of two-pathway experiments based on the conceptual framework of STC model. Two-pathway experiments utilizing either a ‘strong-before-weak’ or ‘weak-before-strong’ protocols, in which strong and weak activity are given sequentially to two independent but convergent inputs, have proven effective in addressing heterosynaptic and late-associative mechanisms (Frey and Morris, 1997). We planned similar experiments reasoning that the potentiation induced by D1R agonists is analogous to strong tetanisation, and input specificity in this case can be achieved by selective silencing of the test stimulation in one input during agonist application (Navakkode et al., 2007). Two stimulating electrodes were positioned in the stratum radiatum of the CA1 region to stimulate two independent synaptic inputs converging onto a common population of neurons and a recording electrode was located midway between the two stimulating electrodes to record the field-EPSP responses (Fig. 2a). SKF was bath applied in three, spaced applications. However, during drug application the baseline stimulation was silenced in one of the inputs (S2) for one hour from the point of drug application. Thus agonist-induced potentiation would be induced only in the active input (S1) but not in the silenced input (S2). After one hour, the test stimulation was resumed in the silenced input and upon 30 minutes of baseline recording, early-LTP (E-LTP) was induced by weak tetanisation (WTET) of S2.

Twenty five micromolars of SKF-induced potentiation in input S1 became significantly different from its baseline after 20 minutes and maintained for 4 hours (Fig. 2b. $P < 0.05$, $n = 7$). However, input S2 in which the baseline stimulation was silenced for one hour showed no potentiation after resuming the recording ($P > 0.05$). Weak tetanisation of S2 resulted in significant potentiation ($P < 0.05$). Interestingly, the WTET, which normally induces only an E-LTP (data not shown), resulted in a late-LTP with a mean potentiation of $(139.98 \pm 10.72)\%$ after two hours that was significantly different from its baseline (Fig. 2b. $P < 0.05$, $n = 7$).

With 10 μM SKF application, input S1 showed significant potentiation by 20 minutes ($P < 0.05$) that sustained for 4 hours (Fig. 2c. $P < 0.05$, $n = 7$). Similar to the experiment with 25 μM SKF, the silenced input S2 did not show potentiation after resuming the baseline stimulation ($P > 0.05$). WTET in S2 resulted in significant potentiation from the baseline ($P < 0.05$). Strikingly, the WTET-induced potentiation did not get reinforced into late-LTP as with 25 μM SKF, but rather decayed to baseline gradually with the mean potentiation of $(108.69 \pm 8.89)\%$ after two hours that was not significantly different from its baseline (Fig. 2c. $P > 0.05$, $n = 7$).

The result was much more intriguing with 5 μM SKF wherein input S1 failed to show significant potentiation [mean potentiation after 100 minutes $(109.78 \pm 7.26)\%$, $P > 0.05$; Fig. 2d, $n = 9$)]. Input S2, upon WTET, showed a significant potentiation ($P < 0.05$, $n = 9$) that rapidly decayed back to baseline becoming not significantly different by 160 minutes ($P > 0.05$).

Late-Plasticity Within the Vicinity of Competing Synapses Prevents Synaptic Competition

From the previous series of experiments, we hypothesized that the decay of both the E-LTP and D1R-mediated potentiation in 5 μM SKF experiments could be because of the competition between these two set of active synapses for a limited pool of PRPs. So, we reasoned that providing additional PRPs by means of strong stimulation within the temporal vicinity of weakly activated synapses should prevent decaying forms of LTP because synthesis of PRPs during late-LTP requires D1R activation (Frey and Schroeder, 1990; Sajikumar and Frey, 2004). For this we designed the three-pathway experiments with three independent but convergent inputs S1, S2, and S3 in the stratum radiatum of CA1 region and a recording electrode positioned in between to record the fEPSP responses (Fig. 3a). Similar to two-pathway experiments, before the application of SKF, the test stimulations in S2 and S3 inputs were silenced for the next one hour, so as to achieve inputspecific D1R-mediated potentiation in S1. After one hour, the
recordings in S2 and S3 were resumed and a stable baseline of 30 minutes was recorded. E-LTP was induced with a WTET in S2 and 15 minutes later late-LTP was induced in S3 by a strong tetanization protocol (STET).

The D1R-LTP in S1 followed the similar time-course as in two-pathway experiments, and the potentiation was not significantly different from the baseline until 90 minutes ($P > 0.05$, $n = 7$). Inputs S2 and S3 showed no significant deviation from baseline upon resuming the stimulation ($P > 0.05$). WTET in S2 resulted in significant potentiation ($P < 0.05$). Significant potentiation was observed in S3 upon STET ($P < 0.05$, $n = 7$) that lasted till the end of the recording [mean potentiation at the end of 4 hours (149.25 ± 7.76%), $P < 0.05$]. Interestingly, the otherwise decaying potentiation was rescued in S1 [mean potentiation at 4 h (119.35 ± 4.13%), $P < 0.05$] and S2 [mean potentiation at 4 h (119.95 ± 2.92%), $P < 0.05$] clearly demonstrating the rescue because of the provision of extra PRPs from S3.

FIGURE 2. Synaptic co-operation and competition mediated by different concentrations of the D1/D5 receptor agonist SKF38393. (a) Schematic representation of a hippocampal slice showing the location of electrodes in CA1 region for the 2-input synaptic tagging and capture experiments. S1 and S2 are two stimulating electrodes in the stratum radiatum that stimulate two independent synaptic inputs to the same neuronal population; one recording electrode is positioned in between the two stimulating electrodes to record field-EPSP from the apical dendrites. S1 and S2 are two independent but convergent inputs. The test stimulation in S2 is silenced just before SKF application for one hour. After resumption and 30 minutes of baseline recording, early-LTP is induced in S2. (b) Twenty five micromolars of SKF induced a slow-onset potentiation in S1 (closed circles) and also transformed the transient form of LTP (early-LTP) induced in S2 (open triangles) into a persistent potentiation ($n = 7$). (c) Ten micromolars of SKF induced a slow-onset potentiation in S1 (closed circles) but the weak tetanization in S2 (open triangles) resulted in early-LTP that was not reinforced ($n = 7$). (d) Five micromolars of SKF-induced slow-onset potentiation in S1 (closed circles) failed to maintain after weak tetanization of S2 (open triangles) and both the responses decayed back to baseline within two hours of early-LTP induction ($n = 9$). Insets show representative fEPSP traces for each input recorded at baseline (solid line, −15 min), 120 min (dotted line), and 240 min (hatched line). Single arrow represents the time point of application of WTET for the induction of early-LTP. Symbols and scale bar for traces as in Figure 1.
Sustained Activation of Extracellular Signal-Regulated Kinases-1 and 2 (ERK1/2) Mediates Differential Effects of D1/D5 Receptor-Dependent Plasticity

Application of dopamine to hippocampal slices has been shown to result in robust activation of p42 MAP kinase in CA1 region (Roberson et al., 1999). Sustained ERK phosphorylation has been reported in the medial prefrontal cortex of 6-hydroxydopamine-lesioned rats following repeated D1R agonist administration which was proposed to play a pivotal role in permanent adaptive changes (Papadeas et al., 2004). Here we investigated whether inhibition of the mitogen-activated protein kinases (MAPKs) ERK1/2 interferes with the maintenance of repeated D1R agonist-induced potentiation. We addressed this by using the specific inhibitors of the upstream kinases of ERK1/2 (MEKs), U0126, and PD98059. The inhibitor was applied one hour after the SKF application and kept throughout until the end of the recording so as to investigate the effect on the LTP maintenance mechanism.

The potentiation induced by 5 μM SKF reached significance on a similar time-course as in the previous experiments.

**FIGURE 3.** Late-plasticity within the vicinity of competing synapses prevents synaptic competition. (a) Schematic representation of a hippocampal slice showing the location of electrodes in CA1 region for the 3-input, synaptic tagging and capture experiments. S1, S2, and S3 were three independent but convergent inputs. The test stimulation in S2 and S3 was silenced just before SKF application for one hour. After resumption and 30 minutes of baseline recording, early-LTP induced in S2 followed, after 15 minutes, by late-LTP induction in S3. (b) A schematic of a pyramidal neuron showing three forms of activity induced in three independent synaptic inputs S1, S2 and S3; 5 μM SKF stimulation in S1, weak tetanization (WTET) in S2, and repeated strong-tetanization (STET) in S3. (c) The otherwise decaying forms of LTP (see Fig. 2d) in S1 (closed circles) and S2 (open triangles) were rescued by the strong-tetanization-induced late-LTP in S3 (closed inverted triangles); (n = 7). Triplet of arrows represent the point of application of STET for the induction of late-LTP. Insets show representative fEPSP traces for each input recorded at baseline (solid line, −15 min), 120 min (dotted line), and 240 min (hatched line). Symbols and scale bar for traces as in Figures 1 and 2.
However, with the application of 5 μM U0126 from 60 minutes onwards, the maintenance of potentiation was affected; the potentiation was only marginally significant by 120 minutes \((P < 0.05)\) and gradually decayed back to baseline becoming not significantly different from the baseline 145 minutes onwards \((P > 0.05\), Fig. 4a, \(n = 7\)). The mean potentiation at the end of 4 hours was \((97.03 \pm 5.28\%) (P > 0.05)\). Similar blockade of the LTP maintenance was observed also with the application of 15 μM PD98059; the mean potentiation at the end of 4 hours was \((97.58 \pm 10.46\%) (Supporting Information Fig. 2a. \(P > 0.999, n = 6\)).

Interestingly, the potentiation induced by 10 μM SKF was also disrupted by the application of 5 μM U0126 (Fig. 4b, \(n = 7\)) or 15 μM PD98059 (Supporting Information Fig. 2b, \(n = 6\)) during the maintenance phase. The potentiation followed the usual time-course and became significantly different by 20 minutes \((P < 0.01)\) and stayed significant until 130 minutes with U0126 \((P < 0.05)\) and until 100 minutes with PD98059 \((P < 0.05)\), later gradually decayed back to baseline. The mean values at the end of 4 hours were not significantly different from the baseline in both cases \((P > 0.05)\).

Surprisingly, 5 μM U0126 application showed no significant effect on the maintenance of potentiation induced by 25 μM SKF (Fig. 4c, \(n = 6\)) as did 15 μM PD98059 (Supporting Information Fig. 2c, \(n = 6\)). The potentiation became significantly different from the baseline by around 30 minutes in both cases \((P < 0.05)\) and stayed significant throughout. The mean potentiation at the end of 4 hours was \((147.99 \pm 11.11\%)\) with U0126 treatment \((P < 0.05)\) and \((135.98 \pm 10.61\%)\) with PD98059 treatment \((P < 0.05)\). We gradually increased the concentration of the inhibitors applied and observed a significant disruption of the maintenance phase within the experimental time-frame with 10 μM U0126 (Fig. 4d, \(n = 6\)) and 30 μM PD98059 (Supporting Information Fig. 2d, \(n = 7\)) application. In these experiments, potentiation induced by 25 μM SKF was only significant up to 115 minutes with U0126 treatment (Fig. 4d, \(P < 0.05\)) and up to 125 minutes with PD98059 treatment (Supporting Information Fig. 2d, \(P < 0.05\)) and decayed to baseline thereafter. At the end of 4 hours, the potentiation was not significantly different from the baseline in both groups \((P > 0.05)\).

A two-way repeated measures ANOVA, comparing different SKF concentration groups and U0126-treatment groups with matched values (i.e., time points 60 min and 240 min compared to baseline) stacked into subcolumns (Fig. 4e), revealed a significant variation between time-points \((F(2,92) = 37.50; P < 0.0001)\) and between groups \((F(6,46) = 4.51; P < 0.001)\). Analysis of simple effects between groups with Tukey’s multiple comparisons test showed significant differences between the ‘SKF-treated’ and ‘SKF+U0126 treated’ groups at 240 minutes. Five micromolars of U0126 treatment significantly blocked potentiation by 240 minutes in 5 μM SKF-treated group, as compared with the 5 μM SKF group without U0126 treatment \((P < 0.05)\). The potentiation induced by 10 μM SKF was also brought back to baseline values by 240 minutes with 5 μM U0126 treatment as compared with the potentiation in 10 μM SKF-treated group \((P < 0.001)\). There was no statistically significant difference in the potentiation at 240 minutes between the 25 μM SKF group and the ‘25 μM SKF+5 μM U0126’ group \((P > 0.05)\). Finally, 25 μM SKF-induced potentiation was significantly blocked by 10 μM U0126 treatment, as evident from the statistically significant difference as compared with 25 μM SKF group \((P < 0.0001)\) and also compared with 25 μM SKF group treated with 5 μM U0126 \((P < 0.001)\).

Similar results were also obtained when these experiments were repeated with another MEK inhibitor PD98059 (Supporting Information Fig. 2e). Though in these series of experiments the concentration of PD98059 required to disrupt the LTP maintenance was higher than that of U0126, it was expected as the potency of this inhibitor is low compared to U0126 (Favata et al., 1998; Levenson et al., 2004). Further, application of either of the MEK inhibitor at the concentrations used above had no significant effect on the baseline fEPSP responses (data not shown).

**DISCUSSION**

Our first series of experiments reveal that the D1R agonist-evoked potentiation at the Schaffer collateral inputs to CA1 apical dendrites exhibits concentration-dependence. With the growing appreciation of the importance of heterosynaptic dopaminergic activation in the induction and maintenance of late-LTP, in certain forms of hippocampus-dependent memory and also in certain neuropsychiatric disorders, these results present a significant aspect to consider both in physiological as well as pathological conditions.

Huang and Kandel (1995) had observed that the D1R agonist application results in a potentiation that develops slowly and peaks by 3-4 hours, whereas in our study the development was comparatively faster. This could be attributed to the methodological differences such as extended preincubation periods and to the drug application paradigms; in that study the agonist was applied in a massed fashion continuously for 15 minutes whereas in our study we used a spaced application protocol (Navakkode et al., 2007; Navakkode et al., 2012).

The two-pathway experiments, designed on the synaptic tagging and capture framework, suggest that the concentration-dependence of dopaminergic modulation has significance during the processing of information from multiple inputs at the cellular level. Sufficiently strong dopaminergic activation might have resulted in the upregulation of a battery of PRPs, which could also be captured by an active weak learning tag. Thus the PRPs, upregulated in sufficiently large amounts, could be shared between both the active set of synapses resulting in a stable plasticity and associativity. These results are consistent with our earlier findings that if plasticity proteins and learning tags are available to interact, the synapses will co-operate with each other for forming stable memory traces (Sajikumar and
FIGURE 4.

DOPAMINERGIC REGULATION OF SYNAPTIC ASSOCIATIVITY

Hippocampus
Frey, 2004). The learning rules change when the concentration of the dopamine decreases. For instance, with lower concentration of D1R agonist, such as 10 μM in our study, the early-LTP induced in the overlapping input was not consolidated. This represents a situation where only one input gets selectively benefitted from the PRPs while not sharing it with the other input active in the critical time-window for association; a situation similar to ‘winner-takes-it-all’ scenario reported recently by us and others (Kano and Hashimoto, 2009; Sajikumar et al., 2014). Thus there was no late-associativity or cooperation between these synapses. The intriguing observation with 5 μM D1R agonist application, wherein potentiation in both inputs decayed to baseline, possibly indicates a scenario similar to ‘competitive maintenance’ (Fonseca et al., 2004) where both the activated set of synapses compete for PRPs from a limited pool which affects the LTP-maintenance in both. Thus the amount of PRPs upregulated at this concentration was not enough to be shared between synapses. This was further supported by the three-pathway experiment, where providing additional PRPs with a strong activity in a convergent input was able to rescue the synapses from competition. Thus it reveals two important aspects of the dynamic nature of the synapses involved in associative memory: (1) With limited availability of plasticity products, synapses will still try to establish a stable memory in the face of competition, sometimes in a winner-takes-all fashion, given that the active synaptic tags were set initially. (2) If enough plasticity products are provided within the vicinity of potentiated synapses, all the synapses will switch to a cooperative state to establish a stable long-term memory trace which would have decayed otherwise because of competition. Thus, we have identified a process in switching the synapses between co-operative to competitive states to deal with different needs of neural computation mediated by dopamine.

Provoked by the interesting results from the co-operative and competitive aspect of synapses, we sought to understand the molecular players involved in the associative maintenance. Given the results, it was clear that the central molecular player in such a scenario should be able to integrate the signals from the D1Rs and the NMDA receptors and transduce the signals for the upregulation of PRPs in a concentration-dependent manner. The potential candidates are the ERK1/2, given their well-documented roles in the mammalian associative learning (Atkins et al., 1998), their capability to act as signal integrators (Roberson et al., 1999; Sweatt, 2001; Selcher et al., 2003; Davis and Laroche, 2006), and their ability to both upregulate translation and also translocate to nucleus and activate transcription (Sweatt, 2001). In hippocampus, a strong synergistic interaction between D1R and NMDAR has been reported (Navakkode et al., 2007), which then leads to a significant activation of ERK1/2 pathway (Sarantis et al., 2009). ERKs are involved in gating late-phase mechanisms of LTP and long-term memory in hippocampus (English and Sweatt, 1996; Blum et al., 1999; Roberson et al., 1999). A particularly interesting observation in our study is that a sustained ERK activation was involved in the maintenance of D1R-mediated long-lasting potentiation. We have also examined the role of other PRPs such as calcium/calmodulin-dependent kinases CaMKII and CaMKIV using different concentrations of the inhibitor KN-93 and our preliminary results show no significant role for these two kinases in the maintenance phase of D1R-mediated potentiation (Unpublished results). Several studies have suggested similar sustained ERK activation in response to a variety of stimuli such as repeated serotonin pulses in aplysia neurons (Sharma et al., 2003), LTP-inducing high frequency stimulation (Ahmed and Frey, 2005), BDNF application (Wu et al., 2001) and in response to memory training in novel taste-learning task (Swank and Sweatt, 2001). Such persistent activation can be the result of many possible scenarios as indicated by Wu et al., (2001). Maharana et al. (2013) found that sustained ERK activation observed in KCl-induced depolarization was dependent on BDNF synthesis and its subsequent activity on its receptor. A propagating ERK activation switch facilitated by feedback events has been proposed to form dendritic zones where plasticity is facilitated (Ajay and Bhalla, 2007). Many other positive-feedback loops may connect active ERK to Raf or MEK activity (Smolen et al., 2008). Bistability of MAPKs has been proposed as a plausible mechanism wherein persistent activity of the kinase and consequent upregulation of translation might contribute to maintenance of LTP and LTM (Smolen et al., 2008). PKMzeta, a persistently active atypical protein kinase C isoform, has been shown to be necessary for maintenance of late-LTP, and some forms of LTM (Serrano et al., 2005; Pastalkova et al., 2006; Sacktor, 2012) and MAPK activity promotes translation of PKM-zeta mRNA (Kelly et al., 2007). PKMzeta was reported to be essential for induction and maintenance of dopamine-LTP in CA1 region (Navakkode et al., 2010). So, it is possible that persistent ERK activity could contribute to LTP maintenance by increasing PKM-zeta synthesis.

![FIGURE 4. ERK1/2 inhibition concentration-dependently blocks the maintenance of D1/D5-agonist-induced potentiation. Following 30 minutes of baseline recording, SKF was applied for three 5-minute durations with 5-minute interval between each application. The MEK-inhibitor, U0126, at the specified concentration, was applied starting 1 h after SKF application till the end of 4 h (shown in rectangles on each graph) (a) 5 μM SKF-induced potentiation was disrupted from maintenance upon application of 5 μM U0126 (n = 7). (b) Application of 5 μM U0126 also disrupted the maintenance of the potentiation induced by 10 μM SKF (n = 7). (c) 5 μM U0126 application had no significant effects on the potentiation induced by 25 μM SKF (n = 6). (d) Maintenance of 25 μM SKF-induced potentiation was disrupted by 10 μM U0126 application causing the potentiation to gradually decay back to baseline (n = 6). (e) Histogram of mean fEPSP slope values recorded for SKF and ‘SKF+U0126’ treatment groups after 60 and 240 minutes of SKF application analyzed with two-way ANOVA. Asterisks indicate significant potentiation compared to a respective baseline group (Tukey’s post-test, *P<0.05, **P<0.01; ***P<0.001; ****P<0.0001). Significant differences between groups is shown with hashtags (Tukey’s post-test, *P<0.05, **P<0.01; ***P<0.001; ****P<0.0001). Error bars indicate ±SEM. Symbols and traces as in Figure 1.](image-url)
Further, our results show that the ERK involvement in D1R agonist-induced potentiation also follows a concentration-dependent activation pattern. Much higher concentration of MEK inhibitors was required to completely block the LTP induced by stronger dopaminergic activation as compared with that required to block the LTP induced by weaker dopaminergic stimulation. This suggests that the level of ERK1/2 activation is more with stronger dopaminergic stimulation than with weaker stimulation. In consistent with our findings, similar dose-dependent activation of ERK1/2 by a D1R agonist has been reported earlier in the mouse prefrontal cortex and also in cultured cortical neurons (Nagai et al., 2007). In a Pavlovian fear conditioning study in amygdala, ERK/MAPK activation was shown to be peaking only by 60 minutes after conditioning and a MEK-inhibitor application impaired fear memory consolidation in a dose-dependent manner and rapidly disrupted the LTP in the slices (Schafe et al., 2000).

The correlation between the activation patterns of the ERK1/2 with the level of potentiation observed at different agonist concentrations suggests a dose-sensor or signal integrator role for ERK1/2 in maintaining plasticity and associative plasticity. ERK activation upon synaptic activity has been proposed to occur in the calcium microdomains in the proximity of synaptic NMDA receptors (Hardingham et al., 2001). It is possible that the stronger dopaminergic activation might involve many such microdomains whereas weaker activation might involve fewer. It has been reported earlier that co-activation of NMDARs and D1/D5-receptors leads to the stimulation of ERK and phosphatidylinositol-3-kinase (PI3 kinase) resulting in the dendritic synthesis of many proteins and also initiates the insertion of glutamate receptors to the synaptic surface (Pfeiffer and Huber, 2006). Active ERK has many downstream targets. ERK activation leads to upregulation of dendritic translation by acting through mTOR pathway (Tsokas et al., 2007; Schicknick et al., 2008) and cap-dependent local translation via MAP-kinase-interacting kinase-1 (MNK1) and eIF4E (Banko et al., 2004; Kelleher et al., 2004). ERK can activate transcription factor cyclic-AMP response element-binding protein (CREB) by multiple routes: by activating the Ribosomal S6-Kinase-2 (RSK2) which can then translocate to nucleus or the phospho-ERK itself can translocate to nucleus (Davis et al., 2000) and directly phosphorylate CREB (Impey et al., 1998). CREB is one of the important plasticity proteins necessary for the establishment of associative plasticity (Barco et al., 2002). Through mediation of RSK2 activation, ERK can also activate the transcription factor Elk-1 (Davis et al., 2000; Sweatt, 2001) which can lead to transcription of Serum-response-element (SRE)-dependent genes, important ones among them being c-fos (Janknecht et al., 1993; Xia et al., 1996) and arc (Waltereit et al., 2001). Though some studies have also shown a role for ERK in certain forms of long-term depression (Thiels et al., 2002; Gallagher et al., 2004) it has been suggested that the specificity of the response might be determined by the pattern of the synaptic activity and by the differential involvement of the many small GTPases (Gallagher et al., 2004).

Given the evidence that ERK-mediated effects on LTP maintenance involves its action both on local translation machinery and nuclear transcriptional events (Davis and Laroche, 2006), an interesting possibility is that the extent of contribution from both of these events might have differential outcomes. The exact feedback mechanisms involved in persistent ERK activation in our study, the signal integration mechanisms, and the downstream targets of ERKs involved in the regulation of synaptic cooperation and competition will be the focus of future studies. Also, it will be interesting to examine whether similar effects could be observed in behavioral studies. However, whether similar mechanisms of ERK activation could be observed in mice is warranted as ERK involvement in certain forms of LTP have been shown to differ between rats and mice (Selcher et al., 2003).

Synaptic associative events such as synaptic tagging and capture have gained a lot of supportive evidence recently with studies demonstrating such mechanisms in anesthetized and freely moving animals, both from electrophysiological recordings and behavioral paradigms (Ballarini et al., 2009; Redondo and Morris, 2011; Shires et al., 2012). Behavioral tagging experiments demonstrating the reinforcement of a weak inhibitory avoidance memory into long-lasting one by the open-field exploration paradigm have indicated the involvement of dopaminergic signaling in such behavioral outcomes (Moncada and Viola, 2007). A similar facilitation has also been shown in a spatial memory task (Wang et al., 2010) Very recently, such a mechanism was also reported to operate in retroactive strengthening of emotional memories in humans (Dunsmaoor et al., 2015). Concomitant recruitment of homosynaptic and heterosynaptic modulatory mechanisms can help tune the specificity of the plasticity and enhance the duration of the plastic change (Bailey et al., 2000).

Taken together, our results present an interesting aspect of dopaminergic modulation of associative information processing in hippocampal CA1 neurons that the dopamine-mediated effects exhibit concentration-dependence leading to differential activation of intracellular mechanisms. Such effects can significantly influence the persistence or elimination of plasticity events occurring in specific dendritic compartments and thus convey different learning rules to same neurons. Given that the extracellular dopamine levels are affected by many drugs used to treat neuropsychiatric disorders and also some of the addictive drugs, our results indicate an important aspect to consider in studies aiming to understand and treat these conditions. The concentration-dependent effects of the modulatory transmitter, as demonstrated for dopamine in the present study, might offer additional computational power and fine-tuning for the cellular information processing mechanisms for long-term associative memory.

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