Changes in Synaptic Morphology Accompany Actin Signaling during LTP

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Stabilization of long-term potentiation (LTP) is commonly proposed to involve changes in synaptic morphology and reorganization of the spine cytoskeleton. Here we tested whether, as predicted from this hypothesis, induction of LTP by theta-burst stimulation activates an actin regulatory pathway and alters synapse morphology within the same dendritic spines. TBS increased severalfold the numbers of spines containing phosphorylated (p) p21-activated kinase (PAK) or its downstream target cofilin; the latter regulates actin filament assembly. The PAK/cofilin phosphoproteins were increased at 2 min but not 30 s post-TBS, peaked at 7 min, and then declined. Double immunostaining for the postsynaptic density protein PSD95 revealed that spines with high pPAK or pCofilin levels had larger synapses (+60–70%) with a more normal size frequency distribution than did neighboring spines. Based on these results and simulations of shape changes to synapse-like objects, we propose that theta stimulation markedly increases the probability that a spine will enter a state characterized by a large, ovoid synapse and that this morphology is important for expression and later stabilization of LTP.

Key words: synaptic plasticity; p21-activated kinase; cofilin; dendritic spines; postsynaptic density; phosphorylation

Introduction

Ultrastructural studies indicate that long-term potentiation (LTP) is accompanied by a change in synaptic morphology, but there is considerable uncertainty as to the nature of the adjustment. Several experiments suggest that a simple change in size and shape is involved (Fifkova and Van Harreveld, 1977; Lee et al., 1979, 1980; Lang et al., 2004; Nagerl et al., 2004; Zhou et al., 2004), whereas others point to a perforation or splitting of the synaptic region (Jones and Calverley, 1991; Muller et al., 2000). These results provide a plausible explanation for the increase in synaptic currents that constitutes LTP expression because of growing evidence that AMPA receptor number scales with the size of the synapse (Nusser et al., 1998; Takumi et al., 1999; Racca et al., 2000; Ganeshina et al., 2004). Possibly related to these morphological observations are light microscopic results showing that intense synaptic activity modifies spine actin networks in cultured cells (Fischer et al., 1998; Matus et al., 2000; Okamoto et al., 2004). Specially driven polymerization has been described for the adult hippocampus (Fukazawa et al., 2003) with recent evidence that AMPA receptor number scales with the size of the synapse (Nusser et al., 1998; Takumi et al., 1999; Racca et al., 2000; Ganeshina et al., 2004). Possibly related to these morphological observations are light microscopic results showing that intense synaptic activity modifies spine actin networks in cultured cells (Fischer et al., 1998; Matus et al., 2000; Okamoto et al., 2004). Specially driven polymerization has been described for the adult hippocampus (Fukazawa et al., 2003) with recent work, using an in situ F-actin labeling procedure, demonstrating the effect within individual spine heads (Lin et al., 2005). Extensions to the latter work indicate that polymerization is essential to the stabilization, rather than the expression, of LTP (Kramar et al., 2006).

The above findings suggest that the theta-burst stimulation (TBS) used to induce synaptic potentiation initiates changes in synaptic anatomy while triggering reorganization of the spine cytoskeleton, with these effects respectively providing for expression and stabilization of LTP. Although this hypothesis accounts for a diverse array of findings, there have been no tests of the essential prediction that TBS rapidly alters actin dynamics and produces a change in synapse morphology within the same spine. The present studies addressed this issue. We first asked whether TBS increases phosphorylation of p21-activated kinase (PAK) and its downstream effector cofilin in spines within adult hippocampal slices. Phosphorylation of cofilin is one of the routes whereby Rho GTPases regulate actin filament assembly (Juliano et al., 2004; Carlisle and Kennedy, 2005). Moreover, the PAK/cofilin pathway is engaged in various cell systems by two receptor classes, integrins and neurotrophin receptors (Gehler et al., 2004; Juliano et al., 2004; Soule et al., 2006), that play prominent roles in theta-induced actin polymerization (Kramar et al., 2006) and spine morphogenesis (Penzes et al., 2003; Boda et al., 2004; Hayashi et al., 2004; Boda et al., 2006). There are thus multiple reasons for assuming that activation of the PAK/cofilin pathway is part of the sequence of events leading from TBS to cytoskeletal reorganization.

In addition to helping to explain how structural changes are initiated during LTP induction, positive results in the PAK/cofilin experiments could open the way to a test of whether theta-induced changes in actin dynamics within spines are associated with changes in synapse morphology. Here we report results that accord with this prediction but that also call for a probabilistic version of the broad hypothesis (for review, see Lynch and...
Baudry, 1984; Yuste and Bonhoeffer, 2001) that the production of stable LTP involves the generation of a new synaptic architecture.

Materials and Methods

All animal procedures were conducted in accordance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals and with protocols approved by the Institutional Animal Care and Use Committee of the University of California, Irvine.

Electrophysiology. Transverse hippocampal slices (350 μm thick from midseptotemporal region) were prepared from young adult (1–2 months of age) male Sprague Dawley rats (Charles River Laboratories, Wilmington, MA) as previously described (Rex et al., 2005) and maintained in an interface recording chamber (95% O2/5% CO2) perfused (60–70 ml/h, 31 ± 1°C) with artificial CSF (aCSF) containing the following (in mM): 124 NaCl, 3 KCl, 1.25 KH2PO4, 3.4 CaCl2, 2.5 MgSO4, 26 NaHCO3, and 10 dextrose (Rex et al., 2005, 2006). Slices were allowed to equilibrate in the recording chamber for 2–3 h before recordings or other treatments were initiated. Unless otherwise noted, chemicals were purchased from Sigma (St. Louis, MO).

Recordings of field EPSPs (fEPSPs) were collected from CA1b stratum (str.) radiatum using a single glass pipette (5 MΩ, 2 m NaCl). Bipolar stimulation (twisted nichrome wires) was delivered alternately at two sites (CA1a and CA1c) in the apical Schaffer-commissural projections (Kramar et al., 2006). The amplitude of the fEPSPs were set to 30–40% of maximum. Low-frequency baseline stimulation was delivered at 0.05 Hz for 20–30 min as described previously (Rex et al., 2005). Slopes and amplitudes of fEPSPs were measured from digitized files (NacGather 2.0; Theta Burst, Irvine, CA) and normalized to the last 10 min of the baseline period. Synaptic potentiation was induced using TBS (10 bursts of four pulses at 100 Hz separated by 200 ms) (Rex et al., 1998). TBS was delivered to the two stimulation electrodes within 5 s of each other, and slices were collected at time points described below; control slices continued to receive baseline stimulation for an equivalent period.

Bath infusion of 6,7-dinitroquinoxaline-2,3-dione (DNOQ; Tocris Bioscience, Ellisville, MO) or DL-2-amino-5-phosphonoveric acid (APV; Tocris Bioscience) was performed via a second infusion line attached to the main input line. DNOQ (20 mM, stored at −20°C) and APV (25 mM, 4°C) stocks were prepared in dimethylsulfoxide and doubly distilled H2O, respectively, and diluted to working concentrations in aCSF before experimentation.

Immunocytochemistry. The following primary antisera and dilutions were used: anti-PAK3 (1:500; catalog #06-902; Upstate, Charlottesville, VA), anti-PAK1 (1:250; catalog #2602; Cell Signaling Technology, Danvers, MA), anti-phospho-PAK (pS141), which recognizes the conserved phosphorylation site on PAKs 1, 2, and 3 (1:100; catalog #44-940G; Biosource, Camarillo, CA), anti-cofilin (1:250; catalog #ACFL02; Cytoskeleton, Denver, CO), anti-phospho-cofilin (pS3; 1:100; catalog #12866; Abcam, Cambridge, MA), and anti-postynaptic density 95 (PSD95; 1:1000; catalog #MA1-045; Affinity Bioreagents, Golden, CO). The anti-PSD95 is a mouse monoclonal; all other primary antisera were raised in rabbit. For preparation of hippocampal slices not receiving electrical stimulation, slices were prepared as described for electrophysiology and maintained in oxygenated aCSF for 2–3 h. For analysis of immunolabeling after electrophysiological stimulation, slices were fixed in 4% paraformaldehyde in 0.1 M sodium phosphate buffer (PB), pH 7.2, for 12–16 h, cryoprotected in 20% sucrose/PB for 2 h at 4°C, sectioned on a freezing microtome at 20 μm, and then mounted onto Fisher Superfrost Plus slides (Fisher Scientific, Pittsburgh, PA). Sections were incubated (40 h, 4°C) in one or more primary antisera in 0.1% PB containing 4% bovine serum albumin and 0.3% Triton X-100 (PBST). Slides were washed in PB (3 × 10 min), incubated (45 min at room temperature) with Texas Red anti-rabbit IgG or fluorescein anti-mouse IgG (1:200; Vector Laboratories, Burlingame, CA) in PBST, rinsed in PB and coveredlipped with Vectashield (Vector Laboratories).

Control tissue was processed by the same procedures but with exclusion of one, or the other, primary antisera from the initial incubation step; under these conditions, the only labeling observed was associated with the primary antibody included in the initial incubation, thereby demonstrating the species specificity of secondary antisera.

Confocal microscopy and measurement of immunoreactive puncta. Laser-scanning confocal microscopy was performed using the Bio-Rad (Hercules, CA) Radiance 2000 Laser Scanning System using a 60 × 1.4 numerical aperture (NA) Plan Apo objective. Optical sections (1.0 μm) were scanned at 4× zoom in a 512 × 512 pixel format. Image montages covering a 205 × 205 μm area were collected from the zone between the two stimulation electrodes containing potentiated synapses (Shimono et al., 2002). The sample field (3126 μm2) selected for puncta quantification was converted to grayscale, and intensity levels were scaled to values determined for each experiment (Photoshop CS, version 8.0; Adobe Systems, San Jose, CA) to visualize low-intensity labeling. Analysis was conducted blind to sample identity on batches of slices that had been sectioned and stained together. In-house built software (Lin et al., 2005; Kramar et al., 2006) was used to count and determine the area of labeled puncta within the size range of spines and synapses. Briefly, intensity thresholds (8 bits/pixel) were applied to identify punctate, spine-like structures at varying levels of label intensity within a range that reliably counted puncta. Thresholds, in practice, had to be placed well above the level at which diffuse, background fluorescence appeared. Lightly labeled elements were therefore lost to the analysis. As a consequence, numbers of a given anatomical structure (i.e., incidence per unit area) as estimated from immunostained sections are significantly smaller than the values recorded with electron microscopy.

Pixel values for each image were normalized to reduce the impact of background intensity differences across the image, binarized using each intensity threshold, and finally cleaned by erosion and dilation filtering (Jain, 1984). Counts from two serial sections were averaged to produce a representative value for each tissue slice. Identified objects <0.04 and >1.2 μm2 were excluded from analysis. Colocalization of immunostaining for phosphorylated (p) cofilin (pCofilin) and pPAK with that for PSD95 was determined automatically (Matlab; Mathworks, Natick, MA) using center coordinates and radius of each labeled object to determine overlap.

Widefield microscopy and three-dimensional reconstruction. Widefield photomicrographs of immunostained tissue sections were acquired with a Leica DM6000 B microscope (Leica Microsystems, Bannockburn, IL) using a 63 × Plan Apo (1.4 NA) objective. Z-series (0.2 μm step) images were deconvolved by iterative restoration using Velocity 4.0 Restoration software (Improvision, Lexington, MA). Three-dimensional reconstructions of the target field were generated and qualitatively analyzed for overlap between phosphoprotein- (pPAK or pCofilin) and PSD95-immunoreactive elements.

Statistical analyses. Analysis of electrophysiological recordings was performed by repeated-measures ANOVA of the final 5 min of experimental recording. Statistical significance of puncta counts was determined by t test or ANOVA followed by Tukey’s HSD post hoc test. Population counts for these experiments represent numbers of slices. Group puncta size distributions were compared using the Kolmogorov–Smirnov Z test. All measures are reported as group means ± SD in text and as means ± SEM in figures.

Results

Theta-burst stimulation causes time-dependent phosphorylation of PAK

The PAK family is composed of six highly conserved GTPase-driven serine/threonine kinases that regulate actin dynamics through parallel pathways that use cofilin, myosin light chain kinase, and other effectors (Hayashi et al., 2004; Juliano et al., 2004; Boda et al., 2006). Two PAK isoforms (PAK3 and PAK1) have been localized to synapses and implicated in spine morphogenesis in the immature hippocampus (Penzes et al., 2003; Boda et al., 2004), but there are as yet no descriptions of the synaptic localization of PAK in adult brain. Accordingly, sections from hippocampal slices were processed for immunocytochemical localization of PAK3 and PAK1 in CA1 str. radiatum using laser-scanning confocal microscopy. PAK3 immunoreactivity was
densely labeled pPAK-immunoreactive (−IR) puncta was much smaller than the number of total PAK-IR puncta (Fig. 1a,b). Counts made with a computerized system provided estimates of 18.3 ± 2.7 puncta/100 µm² for PAK and 1.6 ± 1.6 puncta/100 µm² for pPAK (means ± SDs).

Figure 2 summarizes the number of pPAK+ spine-like profiles present in tissue collected at various times after the delivery of a single train of TBS to each of two collections of Schaffer-commissural projections innervating the analyzed dendritic field (proximal str. radiatum of field CA1b). Evoked responses recorded from this zone (Fig. 2a) confirmed that theta stimulation caused an immediate and dramatic potentiation [traces from slices collected at 2 min or 7 min post-TBS (Fig. 2a)] that stabilized over the subsequent 15–30 min. A pronounced increase in the number of densely pPAK-IR spine-like puncta was apparent in slices collected 2–7 min post-TBS (4.3 ± 1.7 spine/100 µm² (Fig. 2c) compared with controls [1.5 ± 1.9 (Fig. 2b)]. This effect was delayed in onset, because it was not detected at 30 s post-TBS (1.2 ± 1.0 puncta/100 µm²), and transient, in that slices collected at 15–30 min post-TBS (1.3 ± 1.8) did not have a reliably greater than normal number of pPAK+ puncta (Fig. 2d). The overall effect of TBS was significant (ANOVA; p = 0.024), as were the differences between the 2–7 min group and controls or slices collected at 15–30 min (p < 0.05). The numbers of PSD95-IR structures in the same sample fields (Fig. 2e) were not influenced by TBS (ANOVA; p = 0.37). Similarly, antibodies against total PAK3-labeled equal numbers of puncta within str. radiatum of control (19.1 ± 1.0) and TBS-activated slices (18.3 ± 2.7; two-tailed t test, p = 0.59). These results are consistent with evidence that afferent stimulation increases pPAK in spines of dissociated neurons (Hayashi et al., 2004) and confirm the experimental prediction that LTP induction is associated with a rapid and relatively short-lasting activation of the PAK signaling pathway.

Although the relative effects of TBS were large (Fig. 2), the absolute numbers of synapses involved were small. Serial section electron microscopy indicates that ∼250 synapses/100 µm³ are found in str. radiatum of the adult rat hippocampus (Fiala and Harris, 2001) with >95% of these being located on spines. Because the optical sections used in the present analyses were 1 µm thick, the values described in Figure 2 represent per 100 µm³ values. Thus, the observed increase of 3.2 puncta (control vs 2–7 min) would constitute a shift of only ∼1.3% of the total immunolabeled spine population into a pPAK+ state. (It is noteworthy that the number of immunolabeled spines counted in our analysis is undoubtedly an underestimate of the total numbers of spines within the sample field because antibodies do not achieve full tissue penetration, and many lightly labeled spines will fall

Figure 1. Immunostaining for total and pPAK proteins in stratum radiatum of hippocampal field CA1. a, b, Laser-scanning confocal micrographs show the distributions of immunolabeling for PAK3 (a) and pPAK (b). The CA1 cell body layer is to the right (data not shown) in each micrograph. Note that dense pPAK-IR puncta are much less numerous than the comparably sized structures labeled for total PAK3 immunoreactivity. Scale bar, 10 µm. c, Confocal micrographs illustrate the distributions of PAK3 (red), PSD95 (green), and overlapped immunolabeling (yellow in Merge). PAK3-IR puncta were typically associated with dense labeling for PSD95, with the latter either capping (arrow) or overlapping (arrowhead) the PAK3 immunoreactivity. Scale bar, 5 µm. d, Confocal micrographs show profiles labeled with anti-pPAK (red) and anti-PSD95 (green) and the overlap of the two (yellow in Merge). The majority of pPAK-IR profiles were associated with PSD95 immunoreactivity. Scale bar, 5 µm.
into the intensity range at which visual noise begins to intrude on the images.

The above results demonstrate that TBS markedly increases the number of pPAK
+ profiles in the dendritic zone containing the potentiated synapses (i.e., the proximal str. radiatum) but do not establish the regional specificity of the effect. Accordingly, we counted pPAK + profiles in the str. lacunosum-moleculare (<30 μm from the hippocampal fissure), a lamina that does not receive Schaffer-commissural input. The size of the assayed regions and the method for acquiring the images were the same as used for the potentiated zone; slices were collected 2–7 min post-TBS. Counts of pPAK + puncta were not significantly different for control slices (0.5 ± 0.4 puncta/100 μm²) versus those receiving TBS (0.3 ± 0.1 puncta/100 μm²; two-tailed t test, p = 0.60, n = 6/group), thereby demonstrating that effects of theta stimulation on pPAK + spine-like puncta are lamina specific.

Phosphorylation of cofilin
Cofilin accelerates the removal of monomers from the sharp end of actin filaments. This activity is blocked during phosphorylation by LIM kinase, a PAK effector; PAK activation thus results in suppression of cofilin activity and the creation of circumstances favorable for actin cytoskeletal assembly (Carlisle and Kennedy, 2005). Confocal and electron microscopic studies have shown that cofilin is not associated with presynaptic elements and concentrates in dendritic spines in the adult hippocampus (Racz and Weinberg, 2006). We confirmed this localization. Cofilin immunoreactivity was present in large numbers of spine-like puncta scattered throughout the CA1 apical dendritic field (Fig. 3a); counterstaining with antisera to PSD95 confirmed that the great majority of these structures contain postsynaptic densities. In contrast, very few puncta (0.3 ± 0.3/100 μm²) contained high concentrations of pCofilin immunoreactivity and, as with the total protein, these were usually capped by PSD95 immunostaining. To more fully examine this spatial relationship, we used restorative deconvolution analysis extending farther into the head of the spine.

Hippocampal slices receiving TBS exhibited a dramatic, time-dependent increase in numbers of spine-like structures containing dense concentrations of pCofilin immunoreactivity in the dendritic zone innervated by stimulated afferents (Fig. 3e). A more than sixfold increase in pCofilin + puncta was counted in slices collected from 2 to 7 min post-TBS (1.9 ± 1.5 puncta/100 μm²) compared with control slices receiving only low-frequency stimulation (0.3 ± 0.3; p = 0.008, ANOVA, followed by Tukey’s HSD). As was the case for pPAK, the number of pCofilin + puncta was not significantly different from control values in slices fixed 30 s (0.2 ± 0.1; p = 0.99) or 15–30 min (0.8 ± 1.0; p = 0.67) post-TBS. The overall effect of stimulation was highly significant (ANOVA; p = 0.002). Analysis of double labeling for PSD95 again indicated that TBS did not influence numbers of PSD95 + puncta in the sample field (ANOVA; p = 0.23). Slices processed for localization of total cofilin immunoreactivity at 7 min post-TBS similarly showed no effect of stimulation on numbers of cofilin + puncta (10.2 ± 1.38 puncta/100 μm² vs 11.0 ± 1.68 for 7 min post-TBS vs controls, respectively; p = 0.48). These data confirm in detail the previous conclusion that theta stimulation, after a brief delay, activates the PAK/cofilin pathway within the 5–10 min poststimulation period during which LTP undergoes a rapid phase of consolidation.

The sample sizes for the 2–7 min groups in the individual experiments described above were not sufficient to test whether slices collected at 2 min had increased numbers of spines in which the PAK/cofilin pathway had been engaged. We therefore combined cases for the separate markers (anti-pPAK, anti-pCofilin) of the signaling cascade at each time point by converting their values and those of the controls, to Z-scores using the means and

Figure 2. Effects of theta-burst stimulation on numbers of spines with high concentrations of pPAK. a, TBS was delivered to the apical branch of the Schaffer-commissural projections and field EPSPs were recorded from the proximal apical dendrites of field CA1 in hippocampal slices from adult rats. The illustrated results are from one of two converging afferent populations used in each experiment. The traces are responses collected immediately before delivery of TBS (left side) and from the minute before the slices were collected (right). Graphs summarize the mean (± SEM) initial slope of EPSPs elicited by single stimulation pulses pre-TBS and post-TBS for groups of slices (n = 4/group). Calibration: 0.5 mV, 10 ms. TBS was applied at time 0 (arrow). b, c, Laser-scanning confocal micrographs showing pPAK immunostaining in slices that received either control (cont; b) or theta-burst (c) stimulation. Image intensity values are inverted to more easily observe densely labeled puncta. Note that TBS resulted in an increase in the frequency of immunoreactive spine-like puncta (arrows). Scale bar, 10 μm. d, Bar graph showing the number of pPAK + spines in proximal str. radiatum (the recording site in a) from slices fixed 0.5 min (n = 3), 2–7 min (n = 5), or 15–30 min (n = 5) after the delivery of TBS. Control slices (open bar; n = 7) received baseline stimulation. The overall effect of time after stimulation was significant (p = 0.024, one-way ANOVA) and the 2–7 min group was significantly different from both control (p = 0.021, Tukey’s HSD) and 15–30 min post-TBS groups. e, Theta-burst stimulation did not measurably affect the total number of PSD95-IR spines in the same sections in which it increased numbers of pPAK-IR puncta (p = 0.37).
Figure 3. Effects of theta-burst stimulation on the number of spines with high concentrations of pCofilin. a, Laser confocal micrograph shows immunolabeling for total cofilin in the proximal CA1 str. radiatum. Scale bar, 5 μm. b, Micrographs showing immunostaining for pCofilin, PSD95, and the merged image of the two for the same field of CA1 str. radiatum. The pCofilin immunoreactivity (red) was localized to discrete structures that fell within the size range expected for dendritic spines. These were much less numerous than the comparable profiles (yellow at bottom). Scale bar, 0.25 μm. c, d, e, Three-dimensional reconstruction of immunolabeling for pCofilin (red) and PSD95 (green) structures after wide-field microscopic acquisition at 0.2 μm z-steps followed by restorative deconvolution: the adjacent but distinct localizations of pCofilin and PSD95 immunoreactivities are evident. The images are successive 90° rotations (from top to bottom). Scale bar, 0.5 μm. e, Slices were collected 30 s, 2–7 min, or 15–30 min after the delivery of a single train of TBS to the Schaffer–commissural projections and processed for pCofilin immunostaining. Controls (cont; open bar) received baseline stimulation. The marked increase in pCofilin-IR puncta at 2–7 min was highly significant (***p = 0.008, Tukey’s HSD) relative to controls and 30 s post-TBS groups (n = 8 for cont and 2–7 min, n = 9 for 0.5 and 15–30 min). These results provide a close replication of those obtained with pPAK. f, Spine counts were performed within the various groups (control, 30 s post-TBS, etc.) from the separate pPAK and pCofilin experiments. Z-scores were calculated using the means and SDs for the control groups in each experiment; thus, each spine value was expressed as the difference from the mean of the control slices divided by the SD for the control slices. The planned comparisons of 2 and 7 min versus control was highly significant [ANOVA, p = 0.003; **p = 0.003, and ***p = 0.0009 for 2 (n = 7) and 7 (n = 6) min, respectively].

SDs of the pPAK and pCofilin control groups (Fig. 3f). This resulted in groups of 16 control slices and 7 slices collected 2 min post-TBS. A significant increase in labeled puncta was evident in the latter group (p = 0.003) as well as at 7 min post-TBS (n = 6; p = 0.0009, two-tailed t test). The overall effect of time for the combined groups was also highly significant (p = 0.003, ANOVA). It thus appears that the phosphorylation event begins between 30 and 120 s and continues for up to 420 s after delivery of theta stimulation.

The effects of TBS on pCofilin+ puncta were blocked by application of the selective NMDA receptor antagonist (APV) at 50 μM. The APV infusion was initiated 30 min before stimulation and continued until the slices were collected at 7 min post-TBS. The antagonist blocked LTP (p > 0.8 for baseline vs the period from 2 to 7 min post-TBS; repeated-measures ANOVA) and completely eliminated TBS-induced increases in the number of pCofilin+ puncta (control stimulation, 0.55 ± 0.34 puncta/100 μm², n = 5; 7 min post-TBS in the presence of APV, 0.47 ± 0.43 puncta/100 μm², n = 6; p = 0.90 vs control, two-tailed t test).

Phosphorylation in the PAK/cofilin pathway is associated with enlarged synapses

Confirmation that theta-burst stimulation activates the PAK/cofilin cascade opened the way for tests of whether changes in actin dynamics within individual spines are, as proposed, associated with morphologically distinct synapses. Labeled PSD95+ synapses located atop pPAK or pCofilin-immunopositive puncta appeared larger than neighboring contacts in survey micrographs of the CA1 sampling field at 7 min post-TBS (Fig. 4a).

Quantitative analyses confirmed this impression. As expected from earlier counts, the number of pPAK+ puncta was substantially greater at 2–7 min post-TBS than in control slices; the synapses on these pPAK+ puncta (at 2–7 min post-TBS) had a mean PSD95+ area of 0.55 ± 0.12 μm², whereas, in the same fields, the area value for the much more numerous synapses single labeled for PSD95 immunoreactivity was 0.30 ± 0.05 μm² (p = 0.00007, paired t test for n = 8 slices) (Fig. 4b). Surprisingly, this impressive difference was not unique to the 2–7 min post-TBS group: synapses on the occasional pPAK+ puncta were ~70% larger than their neighbors at all experimental time points (30 s and 15 min post-TBS) and in control (no TBS) slices (p = 0.001 in all
cases). It thus appears that induction of LTP in a subgroup of spines in the sampling region converts those spines into a state that is present, but only rarely, under control conditions.

The same pattern of quantitative results was obtained in a second series of slices that were processed for pCofilin immunostaining (Fig. 4b). The mean number of PSD95+ /pCofilin+ puncta was approximately fivefold greater at 2–7 min post-TBS than in control tissue. The PSD95+ synapses associated with pCofilin immunostaining were substantially larger than was the case for synapses that were PSD95+ alone (0.44 ± 0.06 μm² vs. 0.28 ± 0.03 μm², respectively; \( p = 0.00001 \), paired t test for eight slices). It was also the case that synapses associated with the occasional pCofilin+ puncta found at 30 s and 15–30 min post-TBS, and in control slices, were atypically large (data not shown).

Frequency distributions of sizes were generated for PSD95+ synapses that were, or were not, colocalized with pPAK immunoreactivity at 2–7 min post-TBS (Fig. 5a). The synapse frequencies are from the sampling zones of eight slices (18,000 synapses total) and are expressed as a fraction of the total population (pPAK positive or negative) for these slices; the labeled synapses associated with pPAK+ puncta represented 11% of the total population of PSD95+ structures. Sizes for PSD95+ single-labeled synapses formed a Poisson distribution, whereas those for the double-labeled (PSD95+ /pPAK+ ) synapses distributed themselves in a more Gaussian manner; the difference in the frequency distributions was highly significant (Kolmogorov–Smirnov test for eight slices). It

**Figure 4.** Activation of the pPAK/cofilin pathway is associated with large synapses. Slices were collected 2–7 min after delivery of TBS to their Schaffer-commissural afferents. **a,** Micrographs show immunostaining for pCofilin (red), PSD95 (green), and the overlay image (bottom) for a slice harvested 2 min post-TBS. As shown, densely pCofilin-IR puncta (arrows, top) are associated with relatively large PSD95-IR puncta (middle). Scale bar, 1 μm. **b,** Bar graph summarizes the measured areas of PSD95-IR puncta that were double labeled (black bars) for pPAK immunoreactivity (left) or pCofilin immunoreactivity (right) compared with areas of elements labeled for PSD95 alone (open bars) in the same tissue sections. The differences between sizes of PSD95-IR synapses double labeled for the phosphoproteins or labeled for PSD95 immunoreactivity alone was highly significant (**p < 0.01, ***p < 0.001, paired t test vs control; n = 8 slices for both pPAK and pCofilin).

**Figure 5.** Size distributions differ for synapses located on pPAK- and pCofilin-positive spines compared with synapses that are not associated with a phosphoprotein labeled spine. **a,** Frequency distributions for areas of PSD95-IR synapses with (w/) or without (w/o) pPAK immunoreactivity. **b,** TBS had no detectable effects on the size distributions for postsynaptic densities that were not situated on pPAK-IR spines. **c,** Frequency distributions for areas of PSD95-IR synapses that were (closed circles) or were not (open circles) located on pCofilin-positive spines in slices collected 2–7 min post-TBS. Note that the results for this experiment replicate those obtained in the pPAK study (**d**). A comparison of the areas of pCofilin-IR puncta that were (dotted line) or were not (solid line) associated with PSD95 immunostaining. Unlike the reverse case (the sizes of PSD95-IR structures associated with pCofilin shown in **c**), the areas of pCofilin-IR elements were not affected by the presence or absence of double labeling. **e,** There was no correlation within the group of double-labeled structures between synapse size (PSD95-IR profiles) and the area occupied by pCofilin staining (\( r^2 = 0.03 \)).
tested whether double labeling affected the size (and size distribution) of pPAK- and pCofilin-immunostained elements in the slices used in the above analyses (Fig. 5d). The area of pCofilin staining was the same in spine-like profiles with and without associated PSD95 immunoreactivity, whereas the frequency distributions of pCofilin- elements were not detectably different for the single- and double-labeled groups. Thus, double labeling by itself does not change the measured size of target structures. As expected from this conclusion, there was no correlation between the areas of PSD95 + and pCofilin + elements across the population of double-labeled profiles for the 2–7 min groups (r² = 0.03) (Fig. 5e).

In all, then, activation of the PAK/cofilin pathway is associated with a 60–80% increase in the size of postsynaptic densities and a change in the frequency distribution of their areas. A possible explanation for these two effects is described below.

**Simulation of synaptic changes associated with LTP**

A difference in the frequency distribution for sizes, as is evident for the PSD95 + synapses on pPAK- or pCofilin-positive puncta versus those not situated on phosphoprotein labeled spines, is suggestive of a difference in the shape of the objects being sampled. We explored this idea with a simulation in which flattened (i.e., z-dimension is short), ellipsoid discs were placed at random orientations to the vertical in a three-dimensional layer and then projected from a vertical viewpoint onto a flat surface. The shapes were developed from the standard ellipsoid, which is, in canonical form, as follows:

\[
x^2/a^2 + y^2/b^2 + z^2/c^2 = 1,
\]

where a, b, and c are the semi-axes of the ellipsoid. For the purpose of evaluating points on the ellipsoid, it is convenient to use a parametric form of Equation 1, which can be given by the mapping \(\mathbf{f}(\theta, \phi) : \mathbb{R}^2 \rightarrow \mathbb{R}^3\), where

\[
\mathbf{f}(\theta, \phi) = \begin{cases} 
  x(\theta, \phi) \\
  y(\theta, \phi) \\
  z(\theta, \phi)
\end{cases} = \begin{cases} 
  a \cos(\theta) \sin(\phi) \\
  b \sin(\theta) \sin(\phi) \\
  c \cos(\phi)
\end{cases},
\]

0 ≤ \theta ≤ 2π, and 0 ≤ \phi ≤ \pi.

To model the curved nature of the synapse, a parabolic function of x and y were added to the z component of Equation 2, as follows:

\[
z^* = d((\cos(\theta) \sin(\phi))^2 + (\sin(\theta) \sin(\phi))^2) + c \cos(\phi).
\]

Substituting this expression in for the z term in Equation 2 gives the function \(\mathbf{g}(\theta, \phi)\) where

\[
\mathbf{g}(\theta, \phi) = \begin{cases} 
  x(\theta, \phi) \\
  y(\theta, \phi) \\
  z^*(\theta, \phi)
\end{cases}
\]

with \(z^*\) coming from Equation 3. A typical model of a synapse generated from Equation 4 is shown in Figure 6a.

To perform the simulation, the axes values were taken from Gaussian distributions centered at specified means, where the z dimension is held small to achieve shapes that have a flattened appearance. Next, to orient the shapes in space, the rotational transformations about the x-, y-, and z-axes were used:

\[
R_x(\alpha) = \begin{bmatrix} 1 & 0 & 0 \\
0 & \cos(\alpha) & \sin(\alpha) \\
0 & -\sin(\alpha) & \cos(\alpha)
\end{bmatrix},
\]

\[
R_y(\beta) = \begin{bmatrix} \cos(\beta) & 0 & -\sin(\beta) \\
0 & 1 & 0 \\
\sin(\beta) & 0 & \cos(\beta)
\end{bmatrix},
\]

\[
R_z(\gamma) = \begin{bmatrix} \cos(\gamma) & \sin(\gamma) & 0 \\
-\sin(\gamma) & \cos(\gamma) & 0 \\
0 & 0 & 1
\end{bmatrix},
\]

where the angles \(\alpha\), \(\beta\), and \(\gamma\) are chosen from uniformly distributed random variables. Points \(\mathbf{p} \in \mathbb{R}^3\) on a randomly oriented synapse-like shape can then be generated from the following:

\[
\mathbf{p}(\theta, \phi) = R_x(\alpha) R_y(\beta) R_z(\gamma) \mathbf{g}(\theta, \phi) = R_x(\alpha) R_y(\beta) R_z(\gamma) \begin{bmatrix} x(\theta, \phi) \\
y(\theta, \phi) \\
z^*(\theta, \phi)
\end{bmatrix}.
\]

The locus of points generated by projections from Equation 6 onto the x-y plane is bounded by the curve given by the normal to the object when the y-component of the normal is zero. However, because evaluating this curve numerically or obtaining a closed form expression for the curve from Equation 6 is quite complex, a simpler but computationally more expensive approach was taken. In this, a dense series of points were evaluated from applying Equation 6 to the simulated ellipsoid, and these were projected onto a grid in the x-y plane. Elements on this grid behave like pixels on a video monitor in the sense that if enough points are generated on the object, the set of elements in the grid will fill in, and the area of the projected shape can be obtained by simply counting up the filled-in elements.

The areas of the projected shapes, calculated by the means just described, were then measured and plotted as frequency versus size. The projected images are distributed in a Poisson manner.
Figure 7. Effects of an AMPA receptor antagonist on numbers of pCofilin-positive spines. Control and experimental slices were infused with normal aCSF or the AMPA receptor antagonist DNQX, respectively, for 55 min while baseline stimulation was delivered to the Schaffer-commissural fibers. a, Representative traces of fEPSPs collected 10 min before (~10 min) and 40 min after initiation of control or DNQX infusion. Calibration: 0.5 mV, 10 ms. b, Plot of fEPSP slopes for control slices (open circles) and experimental slices treated with DNQX (filled circles) during the interval indicated by the black bar. As shown in a and b, DNQX infusion eliminated Schaffer-commissural fEPSPs. c, Micrographs show pCofilin immunostaining in CA1 str. radiatum of a representative control slice (top) and a DNQX-treated slice harvested ~30 min after the loss of field potentials (bottom). DNQX treatment markedly reduced the number of densely pCofilin-IR structures, although the remaining immunostaining was not qualitatively different from that found in control slices. Scale bar, 5 µm.

**Number of pCofilin-positive puncta in control slices is dependent on excitatory transmission**

The occasional presence of pPAK + and pCofilin + puncta with large synapses in control slices points to the possibility that this combination represents a spine “state” that occurs spontaneously in the absence of high-frequency afferent activity. Such events would necessarily be rare with the state having a half-life of a few minutes; otherwise, the labeled profiles would be expected to be much more common. The question then arises as to whether activation of the PAK/cofilin pathway is a response to randomly occurring combinations of signals impinging on the spine. Perhaps the most likely candidate for a trigger of this kind would be constitutive neurotransmitter release from axon terminals. We tested the idea that glutamatergic transmission might be critical for this phenomenon by incubating slices for 55 min with 10 µM DNQX, an antagonist of AMPA-type glutamate receptors. Stimulation and recording arrangements were as in previous experiments; measures of evoked responses from str. radiatum (Fig. 7a,b) indicate that transmission was blocked for >30 min before fixing the slice. It was evident from surface micrographs (Fig. 7c) that pCofilin + puncta were much less frequent within the physiological recording zone from DNQX-treated slices than was the case for matched controls. Quantitatively, the AMPA receptor antagonist reduced the number of pCofilin + puncta by >50% (two-tailed paired t test, p = 0.003).

**Discussion**

Past work showed that a train of 10 theta bursts causes a marked increase in the number of spines containing high concentrations of F-actin (Lin et al., 2005; Kramar et al., 2006); this effect is eliminated by latrunculin A (Rex et al., 2007), confirming that the stimulation triggered an intense but localized bout of actin polymerization. The actin effect was completely blocked by neutralizing antibodies against β1 family integrins (Kramar et al., 2006) or by a scavenger of BDNF (Rex et al., 2007). This suggests that theta bursts, which release BDNF (Balckowiec and Katz, 2002; Aicardi et al., 2004), engage integrin/trophic factor machinery of a type used by many cell types to translate extracellular signals into morphological changes and altered gene expression. The potent influence of integrins and the TrkB receptor of BDNF over the cytoskeleton involves the Rho family (Geiger et al., 2002; Brakebusch and Fassler, 2003; DeMali et al., 2003; Gehler et al., 2004; Juliano et al., 2004; Wiesner et al., 2005; Chen et al., 2006; Legate et al., 2006), a group of GTPases that collectively act on nucleation and elongation factors as well as on the PAK/cofilin regulatory pathway (Ridley, 2001). The present studies provide the first evidence that this link between surface receptors and the cytoskeleton is activated in mature dendritic spines by theta-burst stimulation. Single theta trains caused severalfold increases in the numbers of spines containing dense concentrations of pPAK and cofilin in the dendritic zones innervated by the stimulated afferents, but not in the adjacent, more distal, dendritic field. Moreover, the time course for the onset of phosphorylation aligned with that described previously for theta-induced actin polymerization (Kramar et al., 2006). It appears, then, that synaptic activity engages known signaling pathways that are used by integrins and BDNF receptors to alter actin dynamics within adult dendritic spines.

The increase in spines with evidence of PAK/cofilin activation agreed reasonably well with estimates of the number of synapses in which LTP occurs. Recordings from pairs of pyramidal cells in adult hippocampal slices suggest that a minimum of two contacts generates a mean evoked (one cell to one cell) EPSP (Debanne et al., 1996; Pavlidis and Madison, 1999) that is <10% of the amplitude of the composite EPSP typically used in LTP experiments. If we assume that ~25 contacts per neuron are involved in the production of the stable (over time) and robust responses used in the present experiments, and take into account the fact that two converging inputs were used, we can estimate that a minimum of 50 synapses per neuron were activated by TBS. Anatomical work indicates that a typical CA1 neuron has 15,000 spines within the entire str. radiatum (Ishizuka et al., 1995; Megias et al., 2001), suggesting that ~7500 spines (per neuron) are found in the proximal half of the lamina containing the activated synaptic population. However, it is unlikely that the greater part of the dendritic...
tree is present within a 350-μm-thick slice. Assuming that one-third of the tree is retained, an estimated 2500 synapses would be found within the target dendritic zone. Combining these values, we can estimate that LTP in the present studies occurred in ~2% of the spine population on a given neuron. This estimate is close to the calculated percentage (1.3%) of the total spine population (~250 synapses/100 μm3) (Sorra and Harris, 1998; Scheff et al., 2005) that exhibited a theta stimulation-induced increase in PAK phosphorylation; this value is likely to be an underestimate because of incomplete labeling with the immunostaining technique.

The discovery that the induction of LTP is associated with phosphorylation of PAK and cofilin opened the way to a test of the long-standing hypothesis that the potentiation effect involves a change in synaptic morphology (Lynch and Baudry, 1984; Yuste and Bonhoeffer, 2001). Early electron microscopic studies showed that the distribution of synaptic lengths changed after induction of LTP (Lee et al., 1979, 1980), a result interpreted as suggesting that a group of rounder profiles had been added to the population. Subsequent studies found that synapses were larger after induction or became split by intervening spine membrane (Jones and Calverley, 1991; Toni et al., 2001; Ostroff et al., 2002; Harris et al., 2003; Lang et al., 2004). Together these findings argue that LTP induction changes synaptic morphology, although no clear consensus as to the nature of the change has emerged. The shape change (of PSD95-IR synapses) observed in the present studies did not involve a simple expansion (i.e., multiplication of size or staining intensity) because the frequency distribution of synaptic areas for synapses on pPAK- or pCofilin-positive spines was very different from that for synapses not associated with the phosphoproteins. A simulation of randomly oriented synapse-like profiles confirmed the suspicion that the observed distributions and greater size could arise if stretched ellipsoids were converted into ovoids. Notably, the transformations involved in the modeling study did not involve any change in synaptic circumference, a result suggesting that a simple rounding of the structures is sufficient to produce the observed increase in area.

Unexpectedly, the few pCofilin-positive spines found in control (no TBS) slices had unusually large synapses; moreover, the frequency distribution of synapse areas showed the same difference between pCofilin-positive and -negative spines as that found after induction of LTP. Given that phosphorylation is a transient event, it seems inevitable that the increase in pCofilin-positive spines occurred sometime in the few minutes before the moment that the slices were collected and fixed. Phosphorylation, in other words, appears to be a spontaneous event. This conclusion implies that the large synapses found on the pPAK+ or pCofilin+ spines also developed spontaneously and are transient. An alternative hypothesis would be that there is a very small population of spines with large synapses and that the PAK/cofilin pathway is continuously activated in these. However, blocking glutamatergic transmission with DNQX, in otherwise untreated slices, markedly reduced the number of spines with dense concentrations of phosphorylated cofilin and attendant large synapses. These results reinforce the conclusion that activation of the PAK/cofilin pathway occurs spontaneously in slices and is associated with the transient emergence of large, presumably ovoid synapses.

Assuming that larger synapses are indicative of enhanced transmission, the above argument implies that the potentiated state is a stochastic event, with a very low probability obtained under resting conditions and a high probability of occurring in spines that received theta-burst stimulation. This would require a low threshold for potentiation, and there are reasons to think that this is the case. For example, using two inputs to the same dendritic targets, it is possible to generate a measurable degree of LTP with a single theta burst (four EPSPs) by activating the other input 200 ms earlier (Larson et al., 1986). It will be of interest in future studies to determine whether the magnitude of the synapse size change is affected by the amount of theta-burst stimulation used. If the number of bursts, or repetition of trains, simply increases the number of spines with large synapses, but does not influence the size distribution of those synapses, it is likely to be the case that LTP is a quantum effect.

The idea that LTP has a low threshold relates only to induction and expression: it would seem that stabilization (consolidation) is another matter. Given the small number of large synapses found in control slices, it must be the case that those developing along with the intermittent, spontaneous activation of the PAK/cofilin pathway quickly revert to a previous, smaller synapse state. It thus appears that the phosphorylation events are not sufficient to trigger the full range of processes that make persistent changes in synaptic morphology. As noted above, actin polymerization and LTP are readily reversed during the first several minutes post-TBS but then become resistant to disruption. The processes responsible for this acquired resistance could thus constitute the high-threshold step in the production of LTP.

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