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Microtubule-associated protein tau is essential for long-term depression in the hippocampus

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The microtubule-associated protein tau is a principal component of neurofibrillary tangles, and has been identified as a key molecule in Alzheimer's disease and other tauopathies. However, it is unknown how a protein that is primarily located in axons is involved in a disease that is believed to have a synaptic origin. To investigate a possible synaptic function of tau, we studied synaptic plasticity in the hippocampus and found a selective deficit in long-term depression (LTD) in tau knockout mice *in vivo* and *in vitro*, an effect that was replicated by RNAi knockdown of tau *in vitro*. We found that the induction of LTD is associated with the glycogen synthase kinase-3-mediated phosphorylation of tau. These observations demonstrate that tau has a critical physiological function in LTD.

1. Introduction

The microtubule-associated protein 'tau' (*MAPT*) gene is located on chromosome 17 and consists of 16 exons [1]. Alternative splicing leads to six isoforms of tau, all of which contain an amino-terminal projection domain and carboxy-terminal with microtubule-binding repeats [2]. Tau contains several critical serine and threonine residues, the phosphorylation of which regulates its binding affinity for microtubules [3,4]. It is believed that through this binding, tau has major roles in stabilizing microtubules [5]. During neuronal development, tau expression is increased in response to nerve growth factor [6], and subsequently enriched in axons, a process that is required for maintaining axon morphology [7]. The extent to which tau may have additional functions unrelated to axonal microtubule stabilization, however, is not known.

Tauopathies, such as Alzheimer's disease (AD), are characterized by widespread accumulation of hyperphosphorylated tau. Once hyperphosphorylated, tau is known to accumulate in somatodendritic compartments and forms the core component of neurofibrillary tangles (NFTs) [8]. It is generally believed that hyperphosphorylation of tau is the critical step in causing it to be missorted from the axon to dendrites, where it interferes with neuronal function [9]. Associated with this accumulation, there is a loss of synapses and eventually neurons [10,11]. However, the mechanism by which this occurs is unknown.

Increasing evidence suggests that in AD, synaptic dysfunction may initiate the cascades that result in cognitive impairment and neurodegeneration. For example, it is well established that oligomeric forms of β -amyloid ($A\beta$) induce a rapid alteration in synaptic plasticity, the process widely believed to underlie learning and memory in the brain [12]. More specifically, $A\beta$ causes inhibition of long-term potentiation (LTP) and enhancement of long-term depression (LTD) in the hippocampus [13]. LTD involves the removal of AMPA receptors (AMPA receptors) from synapses leading to a reduction in synaptic efficiency, and can also result in the shrinkage and elimination of synapses [14]. Therefore, a shift in favour of LTD may lead to neurodegeneration. That such processes may be causally related to neurodegeneration in AD is suggested by the finding that key molecules that are associated with this disorder, such as glycogen synthase kinase (GSK-3 β) and caspase-3, are required for the induction of LTD in the hippocampus [15–18] and mediate the $A\beta$ inhibition of LTP [19]. Interestingly, recent evidence has shown that $A\beta$ inhibition of LTP is absent in the tau knockout (KO) mouse [20]. These data, together with the observation that GSK-3 β directly phosphorylates tau [15,18], suggest that tau may be a downstream effector of GSK-3 β in LTD. Therefore, we decided to examine the role of tau in LTD in the hippocampus.

In this study, we found that in tau KO mice there is a loss of LTD, whereas LTP is not affected. Furthermore, knockdown of tau in hippocampal slices resulted in a complete loss of LTD in the absence of any direct discernible effects on synaptic transmission. We found that LTD was associated with the phosphorylation of tau by GSK-3 β [18]. Collectively, these data suggest that tau phosphorylation is an essential component of LTD.

2. Results

(a) Long-term depression is absent in $MAPT^{+/-}$ and $MAPT^{-/-}$ mice

The physiological role of tau in the hippocampus was initially investigated using tau KO mice. We compared long-term synaptic plasticity in adult (7–11 months old) $MAPT^{+/+}$, $MAPT^{+/-}$ and $MAPT^{-/-}$ mice. Because the tau kinase GSK-3 β is required for LTD in the hippocampus [17], the primary focus of our investigation was on this form of synaptic plasticity. Field excitatory postsynaptic potentials (fEPSPs) were evoked in area CA1 of anaesthetized mice in response to electrical stimulation of the ipsilateral Schaffer collateral–commissural pathway. We found no differences in synaptic transmission between $MAPT^{+/+}$, $MAPT^{+/-}$ and $MAPT^{-/-}$ mice, as assessed using input–output curves (figure 1a), and we observed no significant differences in paired-pulse facilitation over a range of inter-stimulus intervals (figure 1b). However, we found that while

LTD could be readily induced in adult $MAPT^{+/+}$ mice, it was completely absent in $MAPT^{+/-}$ and $MAPT^{-/-}$ mice (figure 1c). By contrast, similar levels of LTP were observed in the three genotypes (figure 1d). Therefore, tau is specifically required for LTD in the hippocampus *in vivo*.

Next, we investigated LTD in acute brain slices from young (14- to 17-day-old) mice. Consistent with the observations *in vivo*, LTD was absent in slices prepared from $MAPT^{-/-}$ mice but was readily induced in slices obtained from $MAPT^{+/+}$ mice (figure 2a). We also investigated LTD induced by a brief application of NMDA (25 μ M, 3 min) and found a specific deficit in slices from the $MAPT^{-/-}$ mice (figure 2b). These results show that the LTD deficit in $MAPT^{-/-}$ mice is apparent early in development and therefore is not directly associated with ageing.

(b) Knockdown of tau by shRNA prevents long-term depression induction

In these experiments, tau was absent or reduced throughout the life of the animals, potentially leading to developmental complications. Therefore, to investigate more directly whether tau is involved in the LTD process, we used an shRNA probe against rat tau and studied synaptic function in rat hippocampal organotypic slice cultures (figure 3). To study the effects of tau knockdown on synaptic transmission, simultaneous recordings of excitatory postsynaptic currents (EPSCs) were performed from tau-shRNA transfected and neighbouring untransfected neurons. There were no significant differences in AMPAR- and NMDA receptor (NMDAR)-mediated EPSCs (EPSC_A and EPSC_N, respectively) between tau-shRNA transfected cells and neighbouring untransfected neurons (EPSC_A in transfected cells, 252 ± 12 pA; EPSC_A in untransfected cells, 253 ± 18 pA, $n = 15$ pairs, $p > 0.05$; EPSC_N in transfected cells, 256 ± 16 pA; EPSC_N in untransfected cells, 268 ± 12 pA, $n = 15$ pairs, $p > 0.05$; figure 3a). We next investigated whether tau-shRNA had any effect on LTD. Consistent with the LTD experiments in $MAPT^{+/-}$ and $MAPT^{-/-}$ mice, tau-shRNA blocked LTD, whereas LTD was routinely induced in simultaneously recorded, neighbouring untransfected cells (tau-shRNA transfected: $92 \pm 3\%$ of baseline; untransfected: $63 \pm 7\%$, $n = 5$, $p < 0.05$, tau-shRNA versus control; figure 3b). The block of LTD was a specific consequence of the knockdown of endogenous tau, because expression of a non-effective, scrambled tau-shRNA had no effect on LTD ($61 \pm 7\%$, $n = 5$, $p > 0.05$, compared with control, figure 3c). Furthermore, the tau-shRNA-mediated LTD deficit was rescued by co-expression of human tau ($60 \pm 4\%$, $n = 5$, $p > 0.05$, compared with control, figure 3d), which was resistant to knockdown (rat tau-shRNA selectively reduces the expression of rat tau but has no effect on human tau; data not shown). These findings are consistent with our KO studies, confirming that tau is required for LTD induction. Taken together, our data show that tau is required for LTD across two species (rats and mice) and in both juvenile and adult tissue.

(c) Tau is found in the postsynaptic compartment

The finding that tau is required for LTD is surprising since LTD is generally considered to be mediated postsynaptically, via the synaptic removal of AMPARs, whereas tau is present primarily in axons. One possibility is that LTD causes the redistribution of tau to dendritic shafts and/or spines. An

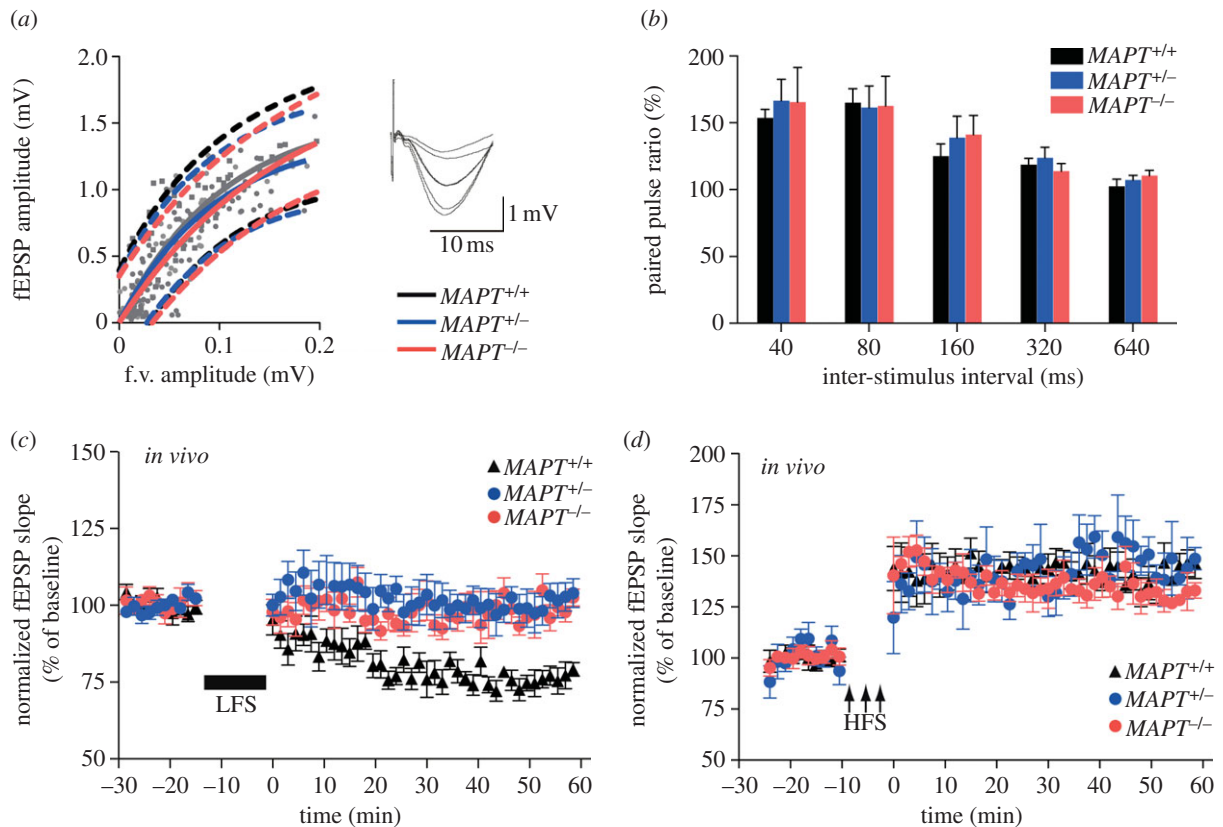


Figure 1. Tau is required for LTD *in vivo* in mice. (a) No differences are shown in synaptic transmission between $MAPT^{+/+}$, $MAPT^{+/-}$ and $MAPT^{-/-}$ mice. The graph plots the fEPSP amplitude versus the fibre volley (f.v.) amplitude (stimulus intensity range: 10–100 μ A, grey dots). Input-output curves show regression (continuous lines) and 95% confidence limits (dashed lines). (b) No differences are shown between $MAPT^{+/+}$, $MAPT^{+/-}$ and $MAPT^{-/-}$ mice in paired-pulse facilitation at various inter-stimulus intervals. (c) LTD is absent in $MAPT^{+/-}$ and $MAPT^{-/-}$ mice. Pooled data from mice (age between 7 and 11 months; $MAPT^{+/+}$: $76 \pm 2\%$; $n = 16$; $MAPT^{+/-}$: $96 \pm 3\%$; $n = 15$; $MAPT^{-/-}$: $98 \pm 2\%$; $n = 11$; $p < 0.001$ in comparison with $MAPT^{+/+}$ mice, Bonferroni's multiple comparison test). (d) No differences are shown in LTP between genotype. Pooled data from 7- to 11-month-old $MAPT^{+/+}$ ($141 \pm 7\%$ of baseline quantified at 60 min after the tetanus, $n = 5$), $MAPT^{+/-}$ ($141 \pm 6\%$, $n = 6$) and $MAPT^{-/-}$ mice ($134 \pm 3\%$, $n = 4$). HFS, high frequency stimulation. (Online version in colour.)

alternative possibility is that a small proportion of tau is normally expressed in a postsynaptic compartment and it is specifically this fraction that is involved in LTD. We explored the latter possibility in two ways. First, we used immunogold electron microscopy (EM) and compared the labelling of tissue from $MAPT^{+/+}$ and $MAPT^{-/-}$ mice (figure 4a). We could detect some immunoreactivity within dendritic spines of the $MAPT^{+/+}$, but not $MAPT^{-/-}$, mice. Second, we probed for the presence of tau, and another microtubule-associated protein (MAP2), in microsomal/organellar (P3), cytoplasmic (S3) and synaptosomal (LP1) fractions prepared from the hippocampus of $MAPT^{+/+}$ mice (figure 4b). As expected, tau and MAP2 were recovered in the P3 fraction and to a lesser extent in the S3 fraction. However, tau was additionally detected in the LP1 fraction. Thus, a proportion of tau is localized at a postsynaptic site where it could, in principle, function directly in LTD.

Because GSK-3 β is a major tau kinase [15,18,21] and is activated during LTD [17], this seemed a likely candidate to mediate the physiological phosphorylation of tau. We hypothesized that the GSK-3 β mediated phosphorylation of tau could be an important regulator of LTD. If this is indeed the case, then a prediction is that the induction of LTD should be associated with an increase in the phosphorylation of tau. To investigate this, we delivered low-frequency stimulation (LFS) and measured the phosphorylation status of tau in the CA1 microdissected dendritic region of rat hippocampal slices (figure 4c). We observed a dramatic increase in the phosphorylation of tau

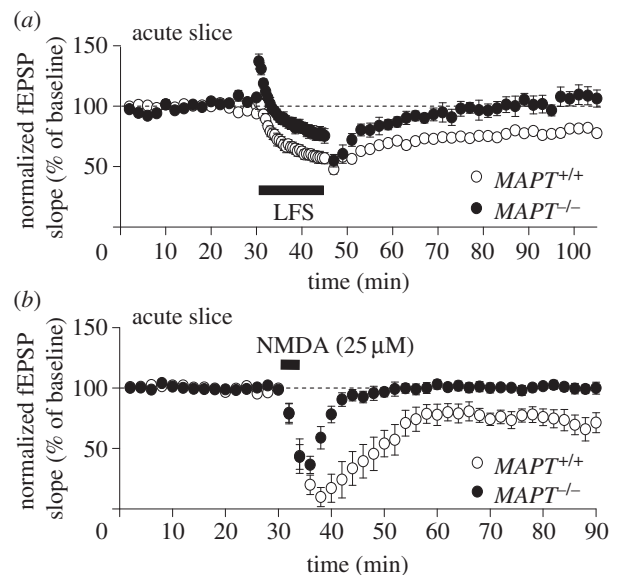


Figure 2. Tau is required for LTD *in vitro* in mice. (a) 1 Hz, 900 pulses induces LTD in $MAPT^{+/+}$ mice ($77 \pm 3\%$, $n = 6$) but is absent in $MAPT^{-/-}$ mice ($106 \pm 7\%$, $n = 5$). Pooled data from postnatal 14- to 17-day-old mice. (b) Bath application of NMDA (25 μ M, 3 min) induces LTD in $MAPT^{+/+}$ mice ($71 \pm 8\%$, $n = 5$) but no LTD in $MAPT^{-/-}$ mice ($99 \pm 4\%$, $n = 6$). Pooled data from postnatal 14- to 17-day-old mice.

using PHF-1 ($p < 0.01$, $n = 4$, figure 4d,e), an antibody that recognizes phosphorylation at residues Ser396 and Ser404 [22], but observed no difference following LFS in the total

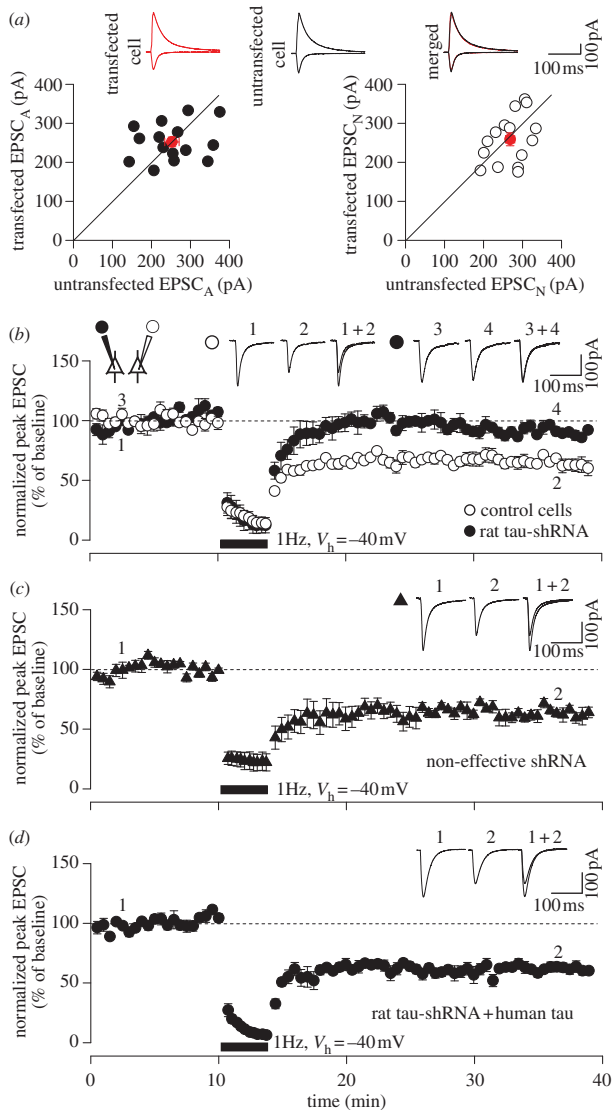


Figure 3. Knockdown of tau blocks LTD *in rats*. (a) Pairwise analysis of basal synaptic transmission between rat tau-shRNA expressing and untransfected neighbouring neurons, each obtained from independent slices. (b) Simultaneous dual-patch recordings were made from rat tau-shRNA-transfected and untransfected neighbouring cells. LFS was delivered (at time indicated by the bar) and LTD quantified 30 min later. (c) Data from scrambled tau-shRNA transfected cells. (d) Data from cells co-expressing rat tau-shRNA and human tau. Error bars indicate s.e.m. (Online version in colour.)

levels of tau using Tau-5 ($p > 0.05$, $n = 4$, figure 4*d,f*), a phosphorylation-independent anti-tau antibody [23]. We next tested whether the phosphorylation of tau was due to activation of GSK-3 during LTD by applying a highly selective GSK-3 β inhibitor, CT-99021, during LFS (figure 4*d*). This treatment eliminated the increase in tau phosphorylation (figure 4*d,e*), while having no effect on the total levels of tau (assessed using Tau-5; figure 4*d,f*). Collectively, these data demonstrate that LFS leads to the phosphorylation of tau in a GSK-3 β -dependent manner, further supporting the idea that this phosphorylation event has a key role in the induction of LTD.

3. Discussion

In this study, we have provided several lines of complementary evidence to suggest that tau is important for LTD in the

hippocampus. First, we have shown that LTD at CA1 synapses *in vivo* is not detectable in mice in which tau is absent or its expression levels are reduced. Second, we found that LTD was absent in slices acutely prepared from juvenile hippocampal tissues of *MAPT*^{-/-} mice. Third, we have demonstrated that knockdown of tau completely blocks LTD in organotypic-cultured slices. Fourth, we have shown that LFS used to elicit physiological LTD leads to enhanced phosphorylation of tau at the PHF-1 epitope, via a GSK-3 β -dependent mechanism.

It is widely believed that tau, under normal conditions, is primarily involved in stabilizing microtubules in axons, and that the dysregulation of this function somehow leads to neuronal pathology [24]. The most prevalent form of such dysregulation occurs through the hyperphosphorylation of tau, which is involved in the generation of NFTs and plays a key role in neurodegenerative conditions such as AD [25]. Hyperphosphorylated tau is missorted to somatodendrites instead of axons, where it is known to accumulate [26–28]. Such missorting is assumed to contribute to neuronal pathology, because it positions tau where it can interfere directly with synaptic function. How tau becomes missorted is not known.

An alternative possibility is that some tau is normally present at synapses and it is this tau that is specifically associated with the neuropathology. Indeed, emerging evidence suggests that tau may be present in dendrites even in the absence of tauopathy [29] and that it could regulate interactions between scaffolding proteins and signalling pathways in the postsynaptic density (PSD) [30]. Furthermore, the localization of tau within the postsynaptic complex can be affected by NMDAR activation [31]. This raises an important question as to what the physiological function of tau in dendrites might be. Here, we have found, using both tau KO mice and RNAi in organotypic slices prepared from rats, that tau is required for LTD. This role is likely to be specific, because we found no evidence that tau is required for maintaining normal synaptic transmission or for LTP. Previous work [9,32] has shown that overexpression of tau may lead to inhibition of LTP. Based on the present findings, we propose that this may be because excess activation of tau induces a chronic form of LTD that is manifest as an impairment in LTP.

We also found that LFS, a physiological LTD induction protocol, resulted in the phosphorylation of tau and that this was dependent on GSK-3. Thus, tau is most probably a physiological substrate of GSK-3 β during LTD. The next key question concerns the physiological downstream effectors of tau during LTD. At present, we can only speculate on this issue. Because tau is a microtubule-associated protein, and because microtubules may be involved in LTD [33], it is possible that tau is involved in the regulation of LTD-dependent microtubule dynamics.

Tau can be divided into a projection domain (towards the N-terminus, encompassing an acidic region and a proline-rich region) and a microtubule-binding domain (towards the C-terminus, including the microtubule-binding repeat region) [34], each having specific roles in the regulation of tau function [35,36]. Within these domains exist multiple regulatory sites of phosphorylation on serine/threonine residues. Our findings suggest a role in LTD for serine residues within the PHF epitope (Ser396/404). Consistent with our findings, Mondragon-Rodriguez *et al.* [31] recently reported a facilitation of Ser396/404 phosphorylation following NMDA treatment. Critically, Ser396/404 residues can both be phosphorylated by GSK-3 β [37,38], an enzyme that is required for the induction

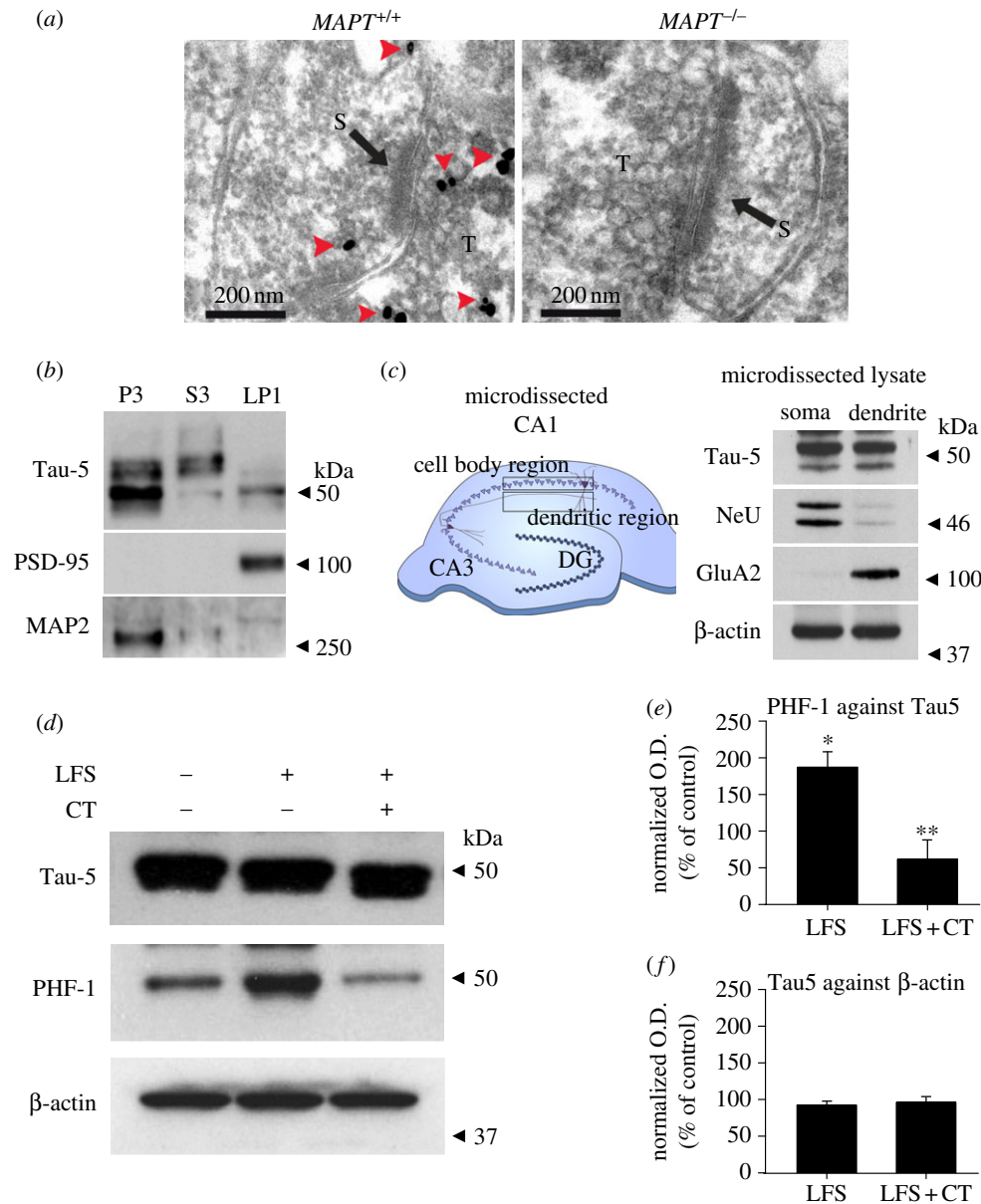


Figure 4. Tau is localized at the synapse and is phosphorylated during LTD. (a) Images of immunoelectron micrographs of hippocampal tissue obtained from *MAPT*^{+/+} (left panel; 4-month-old) and *MAPT*^{-/-} (right panel; 4-month-old) mice. Arrow shows synaptic density and arrowheads indicate tau. JM (rabbit polyclonal anti-tau antibody) and 10 nm gold particle conjugated secondary antibody gave positive signals in *MAPT*^{+/+} but not in *MAPT*^{-/-} mouse tissue. (b) Hippocampus of *MAPT*^{+/+} mouse (4 months old) was fractionated into a microsome/organelle fraction (P3), a cytoplasmic fraction (S3) and a PSD-95-rich fraction (LP1). MAP2 was mostly distributed in the P3 fraction. In comparison, tau (detected using Tau-5) was present in all fractions. (c) Schematic diagram of the microdissection procedure to separate the rat P24–28 CA1 somatic and dendritic regions. Western blotting shows strong expression of NeuN in the somatic region and of GluA2 in the dendritic region. Tau-5 blotting shows expression of tau in both the somatic and dendritic regions. (d) LFS causes an increase in phosphorylation of tau on Ser396/404 (PHF-1 epitope); this LFS-induced increase in phosphorylation of the PHF-1 epitope is attenuated by co-treatment with CT-99021 (CT; 1 μ M). (e) Quantification of PHF-1 levels normalized to Tau-5 in the presence and absence of CT-99021 (control (CTR) versus LFS, * p < 0.05; LFS versus LFS + CT, ** p < 0.01). (f) Quantification of Tau-5 expression, normalized to beta-actin, in the presence and absence of CT-99021 (CTR versus LFS, p > 0.05; LFS versus LFS + CT, p > 0.05). Mann–Whitney non-parametric test was performed to identify changes in statistical significance. (Online version in colour.)

of LTD [17]. However, it was unknown which, if any, of the potential GSK-3 β phosphorylation sites on tau are phosphorylated during the physiological activation of this kinase. Our finding that phosphorylation of Ser396/404 following LFS is prevented by CT-99021 demonstrates that GSK-3 β is upstream of tau in LTD and that this particular phosphorylation event probably has a physiological function.

4. Conclusion

We have shown that tau is required for NMDAR-dependent LTD in the hippocampus. Our data suggest a model whereby

during LTD, activation of GSK-3 β leads to phosphorylation of tau and this promotes LTD.

5. Methods

(a) *In vivo* electrophysiology

Male C57/BL6J mice were used for all comparative KO experiments. *MAPT*^{-/-} and *MAPT*^{+/-} mice were maintained by backcrossing with C57/BL6J mice. Mice were individually housed and kept on a 12 h light/dark schedule. All mice had free access to food and water. Each mouse was anaesthetized with 3% isoflurane–air mixture, and fixed in a stereotaxic device (model 900, David Kopf

Instruments, USA). After exposing the skull, a bipolar-stimulating electrode (two enamel-coated wires with 10 μm diameter and 200 k Ω impedance) was positioned into stratum radiatum of the left hippocampal CA1 area (-1.7 mm from bregma, 1.65 mm from medial, 1.3 mm depth) and a mono-polar recording electrode (0.5–1 M Ω) was placed 200 μm posterior to the stimulating electrode. The animal was maintained anaesthetized (1.5% isoflurane–air mixture) for at least 3 h (body temperature was kept at 36°C). For the fEPSP measurements, the electrical signal was amplified 100 times (ER-1, Cygnus Technology, USA), digitized (Digidata 1321A, Axon Instruments, Foster City, CA, USA) and processed on a computer. To induce LTP, 100 pulses at 100 Hz were applied three times (at 180 s intervals), and to induce LTD, 900 pulses at 1 Hz were delivered. The amplitude and slope of each recorded fEPSP was measured by a custom application based on MATLAB (version 8, Mathworks Inc., CA, USA). fEPSPs were analysed only when the maximal amplitude was over 1 mV, and the latency of the minimum peak from stimulus was shorter than 7 ms. Experiments were performed blindly.

(b) *In vitro* electrophysiology

For *in vitro* electrophysiology experiments, acute hippocampal slices were obtained from P24 to P28 male Wistar rats or *MAPT*^{-/-} and *MAPT*^{+/-} mice. Animals were sacrificed by dislocation of the neck and then decapitated. The brain was rapidly removed and placed in ice-cold artificial cerebrospinal fluid (aCSF) containing (in mM): NaCl 124, KCl 3, NaHCO₃ 26, NaH₂PO₄ 1.25, CaCl₂ 2, MgSO₄ 1 and D-glucose 10 (bubbled with 95% O₂/5% CO₂). Transverse hippocampal slices (400 μm thick) were prepared using a McIlwain tissue chopper (Mickle Laboratory Engineering Co. Ltd., Gomshall, UK). Hippocampal slices were stored in aCSF (20–25°C) for 1–2 h before transferring to the recording chamber, in which they were submerged in aCSF (30°C) flowing at 2 ml min⁻¹. Extracellular field potentials were recorded in the CA1 region using glass electrodes containing NaCl (3 M). A stimulating electrode in CA2 was used to evoke field EPSPs (constant voltage, 100 μs duration, repeated at 30 s intervals). The slope of the evoked fEPSP was measured and expressed relative to the normalized preconditioning baseline. Data were captured and analysed using WinLTP (www.winltp.com). Experiments in which changes in the fibre volley occurred were discarded.

(c) Hippocampal slice culture

Hippocampal slice cultures were prepared from 6- to 7-day-old male Wistar rats, as previously described [19]. Whole-cell patch clamp recordings of CA1 neurons transfected with shRNA plasmids were made 3–4 days following transfection. Using a biolistic Gene Gun (BioRad, USA), neurons were transfected with plasmids expressing shRNA against rat tau protein (OriGene Technologies, MD, USA). A mixture of four different tau (0N/3R, 0N/4R, 1N/4R and 2N/4R) shRNA constructs (1:1:1:1, in pGFP-V-RS vector) were used for tau silencing. A non-effective scrambled sequence shRNA was used as a negative control against tau-shRNA. EPSCs were recorded using a multi-clamp 700B amplifier (Axon Instruments). Recordings were carried out in a solution containing (in mM): NaCl 119, KCl 2.5, CaCl₂ 4, MgCl₂ 4, NaHCO₃ 26, NaH₂PO₄ 1, glucose 11, picrotoxin 0.02 and 2-chloroadenosine 0.01, gassed with 5% CO₂/95% O₂, at pH 7.4. To induce LTD, 200 pulses at 1 Hz were delivered at a holding potential of -40 mV. AMPAR-mediated EPSC amplitude (EPSC_A) was determined as the peak EPSC amplitude at a holding potential of -70 mV. NMDAR-mediated EPSC amplitude (EPSC_N) was determined 50–70 ms after the EPSC_A peak at a holding potential of $+40$ mV. In some experiments, dual patch clamp recordings were made simultaneously from a pair of neighbouring CA1 pyramidal neurons, one transfected and the other untransfected.

n values indicate number of cells, each obtained from independent slices. Error bars indicate s.e.m.

(d) Tau constructs

0N3R and 2N4R human tau cDNAs were framed in pEGFP-C1 host vectors (Clontech, Mountain View, CA, USA), and provided by Drs R. Brandt (University of Ostanbrück, Germany) and S. Lovestone (King's College, UK).

(e) Immunogold electron microscopy

Under deep pentobarbital anaesthesia, animals were perfused with 4% paraformaldehyde in 0.1 M cacodylate buffer (CB, pH 7.4). After further fixation of the brain at 4°C overnight, 300- μm -thick hippocampal slices were made. After incubation with blocking solution (5% normal goat serum in 0.1 M CB) for 1 h at room temperature, the slices were incubated with primary anti-tau antibody, JM (rabbit, 1:300), at 4°C for 2 days, followed by a secondary anti-rabbit IgG conjugated with FITC-gold (goat, Nanoprobes, NY, USA, 1:100) overnight. The slices were re-fixed with a mixture of 2.5% glutaraldehyde and 1% tannic acid at 4°C overnight. The gold signal enhancement procedure was performed according to the manufacturer's instruction (GoldEnhance-EM, Nanoprobes). After the osmication of slices (1% OsO₄–1.5% potassium ferrocyanide in 0.1 M CB) at 4°C for 10 min, the slices were dehydrated, and embedded in epoxy resin. The stratum radiatum of CA1 region was examined electron microscopically (JEM-1200EX, JEOL, Japan) after metal-staining using uranium acetate and lead citrate.

(f) Subcellular fractionation

Partial subcellular fractionation was performed on mouse hippocampus basically according to a previous report [39]. Postnuclear supernatant was subjected to centrifugation (12 500g), and divided into the crude synaptosomal fraction and synaptosome-depleted fraction. The crude synaptosomal fraction was further purified by hypotonic lysis and centrifugation (25 000g), and the resultant pellet was the PSD-95-rich synaptosome fraction (LP1). The synaptosome-depleted fraction was further subjected to ultracentrifugation (100 000g), and separated to the microsome fraction (P3) and cytoplasm fraction (S3). We used the following antibodies for experiments: NeuN, mouse monoclonal (Millipore; 1:1000) Tau-5, mouse monoclonal (Invitrogen; 1:500); PSD-95 (Millipore; 1:1000) and MAP2 (Millipore; 1:1000).

(g) Low-frequency stimulation, microdissection and western blotting

Rat hippocampal slices from P24 to P28 were subjected to a standard 1 Hz, 900 pulses, LFS protocol (stimulation set at a predetermined intensity; 70% of the maximal fEPSP amplitude voltage stimulation), using a bipolar stimulation electrode, in the presence or absence of CT-99021 (1 μM). The dendritic CA1 region was then immediately dissected and snap frozen. Samples were lysed and SDS-PAGE was performed as previously shown [19]. PHF1 monoclonal antibody (kindly provided by Dr Peter Davies) was used at 1:1000 for western blot. Optical densities of immunoreactive bands were quantified using NIH IMAGEJ software (downloaded from <http://rsb.info.nih.gov/ij/>). *n* indicates the number of independent experiments from different animals.

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References

- Andreadis A, Brown WM, Kosik KS. 1992 Structure and novel exons of the human tau gene. *Biochemistry* **31**, 10 626–10 633. (doi:10.1021/bi00158a027)
- Goedert M, Wischik CM, Crowther RA, Walker JE, Klug A. 1988 Cloning and sequencing of the cDNA encoding a core protein of the paired helical filament of Alzheimer disease: identification as the microtubule-associated protein tau. *Proc. Natl Acad. Sci. USA* **85**, 4051–4055. (doi:10.1073/pnas.85.11.4051)
- Lindwall G, Cole RD. 1984 Phosphorylation affects the ability of tau protein to promote microtubule assembly. *J. Biol. Chem.* **259**, 5301–5305.
- Biernat J, Gustke N, Drewes G, Mandelkow EM, Mandelkow E. 1993 Phosphorylation of Ser262 strongly reduces binding of tau to microtubules: distinction between PHF-like immunoreactivity and microtubule binding. *Neuron* **11**, 153–163. (doi:10.1016/0896-6273(93)90279-Z)
- Weingarten MD, Lockwood AH, Hwo SY, Kirschner MW. 1975 A protein factor essential for microtubule assembly. *Proc. Natl Acad. Sci. USA* **72**, 1858–1862. (doi:10.1073/pnas.72.5.1858)
- Drubin DG, Feinstein SC, Shooter EM, Kirschner MW. 1985 Nerve growth factor-induced neurite outgrowth in PC12 cells involves the coordinate induction of microtubule assembly and assembly-promoting factors. *J. Cell Biol.* **101**, 1799–1807. (doi:10.1083/jcb.101.5.1799)
- Caceres A, Kosik KS. 1990 Inhibition of neurite polarity by tau antisense oligonucleotides in primary cerebellar neurons. *Nature* **343**, 461–463. (doi:10.1038/343461a0)
- Kowall NW, Kosik KS. 1987 Axonal disruption and aberrant localization of tau protein characterize the neuropil pathology of Alzheimer's disease. *Ann. Neurol.* **22**, 639–643. (doi:10.1002/ana.410220514)
- Sydow A *et al.* 2011 Tau-induced defects in synaptic plasticity, learning, and memory are reversible in transgenic mice after switching off the toxic Tau mutant. *J. Neurosci.* **31**, 2511–2525. (doi:10.1523/JNEUROSCI.5245-10.2011)
- Kimura T *et al.* 2010 Aggregation of detergent-insoluble tau is involved in neuronal loss but not in synaptic loss. *J. Biol. Chem.* **285**, 38 692–38 699. (doi:10.1074/jbc.M110.136630)
- Kimura T *et al.* 2007 Hyperphosphorylated tau in parahippocampal cortex impairs place learning in aged mice expressing wild-type human tau. *EMBO J.* **26**, 5143–5152. (doi:10.1038/sj.emboj.7601917)
- Bliss TV, Collingridge GL. 1993 A synaptic model of memory: long-term potentiation in the hippocampus. *Nature* **361**, 31–39. (doi:10.1038/361031a0)
- Walsh DM, Klyubin I, Fadeeva JV, Cullen WK, Anwyl R, Wolfe MS, Rowan MJ, Selkoe DJ. 2002 Naturally secreted oligomers of amyloid β protein potently inhibit hippocampal long-term potentiation *in vivo*. *Nature* **416**, 535–539. (doi:10.1038/416535a)
- Collingridge GL, Peineau S, Howland JG, Wang YT. 2010 Long-term depression in the CNS. *Nat. Rev. Neurosci.* **11**, 459–473. (doi:10.1038/nrn2867)
- Lovestone S *et al.* 1994 Alzheimer's disease-like phosphorylation of the microtubule-associated protein tau by glycogen synthase kinase-3 in transfected mammalian cells. *Curr. Biol.* **4**, 1077–1086. (doi:10.1016/S0960-9822(00)00246-3)
- Li Z, Jo J, Jia JM, Lo SC, Whitcomb DJ, Jiao S, Cho K, Sheng M. 2010 Caspase-3 activation via mitochondria is required for long-term depression and AMPA receptor internalization. *Cell* **141**, 859–871. (doi:10.1016/j.cell.2010.03.053)
- Peineau S *et al.* 2007 LTP inhibits LTD in the hippocampus via regulation of GSK3 β . *Neuron* **53**, 703–717. (doi:10.1016/j.neuron.2007.01.029)
- Takashima A, Honda T, Yasutake K, Michel G, Murayama O, Murayama M, Ishiguro K, Yamaguchi H. 1998 Activation of tau protein kinase I/glycogen synthase kinase-3 β by amyloid β peptide (25–35) enhances phosphorylation of tau in hippocampal neurons. *Neurosci. Res.* **31**, 317–323. (doi:10.1016/S0168-0102(98)00061-3)
- Jo J, Whitcomb DJ. 2011 A β (1–42) inhibition of LTP is mediated by a signaling pathway involving caspase-3, Akt1 and GSK-3 β . *Nat. Neurosci.* **14**, 545–547. (doi:10.1038/nn.2785)
- Shipton OA *et al.* 2011 Tau protein is required for amyloid β -induced impairment of hippocampal long-term potentiation. *J. Neurosci.* **31**, 1688–1692. (doi:10.1523/JNEUROSCI.2610-10.2011)
- Cho JH, Johnson GVW. 2003 Glycogen synthase kinase 3 β phosphorylates tau at both primed and unprimed sites. Differential impact on microtubule binding. *J. Biol. Chem.* **278**, 187–193. (doi:10.1074/jbc.M206236200)
- Otvos Jr L, Feiner L, Lang E, Szendrei GI, Goedert M, Lee VM. 1994 Monoclonal antibody PHF-1 recognizes tau protein phosphorylated at serine residues 396 and 404. *J. Neurosci. Res.* **39**, 669–673. (doi:10.1002/jnr.490390607)
- LoPresti P, Szuchet S, Pappasozomenos SC, Zinkowski RP, Binder LI. 1995 Functional implications for the microtubule-associated protein tau: localization in oligodendrocytes. *Proc. Natl Acad. Sci. USA* **92**, 10 369–10 373. (doi:10.1073/pnas.92.22.10369)
- Lee VM, Goedert M, Trojanowski JQ. 2001 Neurodegenerative tauopathies. *Annu. Rev. Neurosci.* **24**, 1121–1159. (doi:10.1146/annurev.neuro.24.1.1121)
- Takashima A. 2010 Tau aggregation is a therapeutic target for Alzheimer's disease. *Curr. Alzheimer Res.* **7**, 665–669. (doi:10.2174/156720510793611600)
- Hoover BR *et al.* 2010 Tau mislocalization to dendritic spines mediates synaptic dysfunction independently of neurodegeneration. *Neuron* **68**, 1067–1081. (doi:10.1016/j.neuron.2010.11.030)
- Iltner LM *et al.* 2010 Dendritic function of tau mediates amyloid- β toxicity in Alzheimer's disease mouse models. *Cell* **142**, 387–397. (doi:10.1016/j.cell.2010.06.036)
- Zempel H, Thies E, Mandelkow E, Mandelkow EM. 2010 A β oligomers cause localized Ca²⁺ elevation, missorting of endogenous Tau into dendrites, Tau phosphorylation, and destruction of microtubules and spines. *J. Neurosci.* **30**, 11 938–11 950. (doi:10.1523/jneurosci.2357-10.2010)
- Iltner LM, Gotz J. 2011 Amyloid- β and tau: a toxic pas de deux in Alzheimer's disease. *Nat. Rev. Neurosci.* **12**, 65–72. (doi:10.1038/nrn2967)
- Morris M, Maeda S, Vossell K, Mucke L. 2011 The many faces of tau. *Neuron* **70**, 410–426. (doi:10.1016/j.neuron.2011.04.009)
- Mondragon-Rodriguez S, Trillaud-Doppia E, Dutilot A, Bourgeois C, Lauzon M, Leclerc N, Boehm J. 2012 Interaction of endogenous tau protein with synaptic proteins is regulated by N-methyl-D-aspartate receptor-dependent tau phosphorylation. *J. Biol. Chem.* **287**, 32 040–32 053. (doi:10.1074/jbc.M112.401240)
- Lasagna-Reeves CA *et al.* 2012 Alzheimer brain-derived tau oligomers propagate pathology from endogenous tau. *Sci. Rep.* **2**, 700. (doi:10.1038/srep00700)
- Hoogenraad CC, Akhmanova A. 2010 Dendritic spine plasticity: new regulatory roles of dynamic microtubules. *Neuroscientist* **16**, 650–661. (doi:10.1177/1073858410386357)
- Buée L, Bussièrre T, Buée-Scherrer V, Delacourte A, Hof PR. 2000 Tau protein isoforms, phosphorylation and role in neurodegenerative disorders. *Brain Res. Brain Res. Rev.* **33**, 95–130.

35. Lee G, Neve RL, Kosik KS. 1989 The microtubule binding domain of tau protein. *Neuron* **2**, 1615–1624. (doi:10.1016/0896-6273(89)90050-0)
36. Brandt R, Leger J, Lee G. 1995 Interaction of tau with the neural plasma membrane mediated by tau's amino-terminal projection domain. *J. Cell Biol.* **131**, 1327–1340. (doi:10.1083/jcb.131.5.1327)
37. Hanger DP, Hughes K, Woodgett JR, Brion JP, Anderton BH. 1992 Glycogen synthase kinase-3 induces Alzheimer's disease-like phosphorylation of tau: generation of paired helical filament epitopes and neuronal localisation of the kinase. *Neurosci. Lett.* **147**, 58–62. (doi:10.1016/0304-3940(92)90774-2)
38. Liu SJ *et al.* 2004 Tau becomes a more favorable substrate for GSK-3 when it is prephosphorylated by PKA in rat brain. *J. Biol. Chem.* **279**, 50 078–50 088. (doi:10.1074/jbc.M406109200)
39. Huttner WB, Schiebler W, Greengard P, De Camilli P. 1983 Synapsin I (protein I), a nerve terminal-specific phosphoprotein. III. Its association with synaptic vesicles studied in a highly purified synaptic vesicle preparation. *J. Cell Biol.* **96**, 1374–1388. (doi:10.1083/jcb.96.5.1374)