Microtubule affinity-regulating kinase 2 functions downstream of the PAR-3/PAR-6/atypical PKC complex in regulating hippocampal neuronal polarity


*Institute of Neuroscience and Key Laboratory of Neurobiology, Shanghai Institute for Biological Sciences, and †Graduate School of Chinese Academy of Sciences, Chinese Academy of Sciences, 320 Yue Yang Road, Shanghai 200031, China; ‡Departments of Cell Biology and Physiology and Internal Medicine, Howard Hughes Medical Institute, Washington University School of Medicine, 660 South Euclid Avenue, St. Louis, MO 63110; and §Cellzome AG, Meyerhofstrasse 1, D-69117 Heidelberg, Germany

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The PAR-3/PAR-6/atypical PKC (aPKC) complex is required for axon–dendrite specification of hippocampal neurons. However, the downstream effectors of this complex are not well defined. In this article, we report a role for microtubule affinity-regulating kinase (MARK)/PAR-1 in axon–dendrite specification. Knocking down MARK2 expression with small interfering RNAs induced formation of multiple axon-like neurites and promoted axon outgrowth. Ectopic expression of MARK2 caused phosphorylation of tau (S262) and led to loss of axons, and this phenotype was rescued by expression of PAR-3, PAR-6, and aPKC. In contrast, the polarity defects caused by an MARK2 mutant (T595A), which is not responsive to aPKC, were not rescued by the PAR-3/PAR-6/aPKC complex. Moreover, polarity was abrogated in neurons overexpressing a mutant of MARK2 with a deleted kinase domain but an intact aPKC-binding domain. Finally, suppression of MARK2 rescued the polarity defects induced by a dominant-negative aPKC mutant. These results suggest that MARK2 is involved in neuronal polarization and functions downstream of the PAR-3/PAR-6/aPKC complex. We propose that aPKC in complex with PAR-3/PAR-6 negatively regulates MARK(s), which in turn causes dephosphorylation of microtubule-associated proteins, such as tau, leading to the assembly of microtubules and elongation of axons.

Results

Suppressing MARK2 Induces Formation of Multiple Axons. MARK2 is negatively regulated by aPKC, which phosphorylates MARK2 on T595 (26). Interestingly, the ratio of p-MARK2 (T595)/MARK2 in axon tips of stage-3 neurons was significantly higher than that in dendrite tips (for details, see Supporting Results and Fig. 6, which are published as supporting information on the PNAS web site). To explore the function of MARK2, we designed two small interfering RNAs (siRNA), siRNA-1677 and -1180. siRNA-1677 suppressed MARK2 expression in transfected human embryonic kidney HEK-293 cells compared with control siRNA, whereas siRNA-1180 had little effect (Fig. 1A). Importantly, expression of EGFp-tau was not affected by the MARK2-specific siRNA (Fig. 1A). The effect of siRNA was also examined in primary neurons in which endogenous MARK2 was suppressed by siRNA-1677 but not siRNA-1180 (Fig. 1B). To examine the effects of the MARK2-

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Abbreviations: MAP, microtubule-associated protein; MT, microtubule; GSK3β, glycogen synthase kinase 3β; aPKC, atypical PKC; PAR, partitioning-defective protein; MARK, microtubule affinity-regulating kinase; siRNA, small interfering RNA; HEK, human embryonic kidney; DIVn, n days in vitro; aPKC-N, amino-terminal PB1 domain of aPKC.

†To whom correspondence should be addressed. E-mail: zgluo@ion.ac.cn.

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siRNA on neuronal polarity, hippocampal neurons were electroporated with MARK2-siRNAs, together with an EGFP plasmid before plating and were analyzed at 3 days in vitro (DIV3) for their ability to polarize. As seen in Fig. 1, neurons expressing EGFP alone showed normal polarity, as indicated by the presence of a single process labeled with the axonal marker Tau1 (Fig. 1C). In contrast, the majority (~63%) of neurons transfected with siRNA-1677 developed multiple long processes that stained positive for Tau1, identifying them as axons (Fig. 1 D and G). Only a small fraction of neurons transfected with either control siRNA (~13%) or siRNAi-1180 (~12%) grew multiple axon-like processes (Fig. 1G). Fig. 7, which is published as supporting information on the PNAS web site, showed that siRNA-1677-treated neurons had multiple long processes labeled with TUJ1, and these processes were longer than the single axons of neighboring nontransfected cells (see Fig. 7C). Synapsin, another axonal marker in mature neurons (24), was also seen in the multiple long processes generated in MARK2-siRNA-treated cells at DIV5 (Fig. 1E), further confirming their axonal properties. Transfection with MARK2-siRNA-1677 caused a 2-fold increase in axon length, whereas MARK2-siRNA-1180 had no apparent effect (Fig. 1F). Interestingly, siRNA-1677-transfected cells often exhibited numerous long protrusions along their axon-like processes (see Fig. 7 D, F, and G), reminiscent of dynamic F-actin (see Fig. 7B). These structures were rarely observed in control cells (see Fig. 7E).

**aPKC Inhibits MARK2 Phosphorylation of Tau.** How does MARK2 specify axon formation? MARK phosphorylates MAPs, such as doublecortin, MAP2, MAP4, and tau, to trigger MT disassembly (21, 22, 25). As shown in Fig. 2, knocking down MARK2 expression with siRNA-1677 in primary neurons resulted in a decrease of tau phosphorylation at S262 and an increase of Tau1, whereas the level of total tau was not altered (Fig. 2A). This result suggests the involvement of MARK2 in phosphorylating tau in vivo. aPKC associates with and phosphorylates MARK2/PAR-1 on a specific site, T595 in PAR-1b (20, 26, 27), leading to a loss of plasma membrane binding and inhibition of MARK2/PAR-1b kinase activity (26). Consistent with the positive staining of p-MARK2 (T595) in neurons at stage 3 (Fig. 6F), immunoblotting revealed that the level of p-MARK2 (T595) in hippocampal neurons was high on day 3 (with most neurons in stage 3) but low on day 1 (with most neurons in stages 1 and 2) (Fig. 2B). Bisindolylmaleimide, an inhibitor of PKC, significantly decreased MARK2 phosphorylation at T595 (Fig. 2B). In agreement with this result, overexpression of MARK2 in HEK-293 cells or PC12 cells increased tau phosphorylation at S262 (Fig. 2C and see Fig. 8A), which is published as supporting information on the PNAS web site). However, when aPKC was cotransfected with MARK2, the phosphorylation of tau (S262) was significantly decreased (Fig. 2C and D and see Fig. 8A). In contrast, aPKC did not block the ability of MARK2T595A, a mutant that retains kinase activity but cannot be phosphorylated by aPKC (26), to phosphorylate tau (Fig. 2 C, lanes 4 and 5, and D).

The SAP kinases contain a kinase domain related to that of MARK2 (50–52% amino acid homology in the kinase domain) but diverge outside of the kinase domain (28). Consistent with ref. 28, overexpression of SAD-B caused phosphorylation of tau (S262) (Fig. 2 E, lanes 1 and 2). However, SAD phosphorylation of tau (S262) did not appear to be regulated by aPKC, even in the presence of PAR-3 and -6 (Fig. 2 E, lanes 2–4). These results demonstrate the specificity of aPKC for MARK2.

**MARK Is Involved in Neuronal Polarity Specification.** MARK2 has a kinase domain at its amino terminus, followed by a divergent middle...
(M) region and a conserved carboxyl-terminal (C) region (Fig. 3A) (29). To test which domain of MARK2 interacts with aPKC, HEK-293 cells were transfected with vectors encoding Flag-aPKC together with EGFP-MARK2 or mutants. As shown in Fig. 3, aPKC associated with MARK2 and the fragment containing the M and C regions (amino acids 321–691), but not the M region alone (amino acids 321–582) (Fig. 3B). To further explore the role of MARK2 in neuronal polarity, primary neurons were transfected with wild-type or mutant forms of MARK2. Ectopic expression of MARK2 resulted in enhanced phosphorylation of tau at S262 (Fig. 3C). The level of dephospho-tau (Tau1) was decreased by MARK2, whereas total tau remained constant (Fig. 3C). Interestingly,

Fig. 3. The role of MARK2 and its mutants in neuronal polarity. (A) Schematic diagrams of MARK2 structure. (B) HEK-293 cells were transfected with Flag-aPKC and EGFP-tagged MARK2 constructs. The immunoprecipitates with anti-MARK2 antibody were subjected to immunoblotting with indicated antibodies. IgG-H, IgG heavy chain. (C) Primary neurons were transfected with MARK2 constructs or control plasmid and cultured for 48 h. The levels of different tau isoforms were accessed by immunoblotting. (D) Neurons transfected with EGFP-MARK2 were stained at DIV3 with antibodies against different tau isoforms, Tau1, and p-tau (S262). Note that the EGFP-MARK2-positive cell (arrow) stained strongly for p-tau (S262) but weakly for Tau1, in comparison with the neighboring nontransfected cell (arrowhead). (E) Neurons transfected with EGFP-MARK2MC or EGFP were stained with anti-TUJ1 antibody at DIV3. Note that EGFP-MARK2MC-expressing cells were unpolarized, whereas the neighboring untransfected cells were polarized. (F) Neurons transfected with EGFP-MARK2MC were stained with Tau1 antibody at DIV3. (G) Neurons transfected with EGFP-MARK2MC were stained with Tau1 antibody at DIV7. Note that MARK2MC-expressing cells (arrows) stained low for Tau1 signal in comparison with the neighboring nontransfected cell. (H) Quantitative analysis of neuronal polarity. None, neurons with short neurites in similar length; single, neurons with one Tau1-positive process >100 μm and at least twice as long as the second longest process; multiple, neurons with two or more Tau1-positive processes >100 μm and twice as long as other neurites. The data were obtained from three independent experiments. At least 90 cells for each construct were analyzed per experiment.
MARK2MC, which lacks kinase domain but retains aPKC-binding ability, also caused an increase of \(p\)-tau (S262) and a decrease of Tau1 (Fig. 3C). MARK2M, however, which does not interact with aPKC, had no apparent effect on tau phosphorylation (Fig. 3C).

Next, we characterized the effects of MARK2 on neuronal polarity by expressing wild-type or mutant forms of MARK2 in hippocampal neurons. After being transfected with EGFP-MARK2, only a few neurons attached to poly-D-lysine-coated plates; these neurons showed increased \(p\)-tau (S262) staining, but even after culturing for 72 h, they did not develop axons (Figs. 3D and 4E). We used MARK2 mutants to investigate the role of MARK2/aPKC interactions in neuronal polarity. Interestingly, the majority of neurons expressing EGFP-MARK2MC did not polarize. These neurons possessed several minor processes labeled with TUJ1 and lacked a single axon-like long process (polarized neurons with a single axon, 54 ± 6.1% for MARK2KD and 85 ± 6.1% for EGFP; \(P < 0.01\); Fig. 3H).

PAR-3/PAR-6/aPKC Counteract MARK2 in Axon Formation. Inhibition of MARK2 by aPKC is predicted to decrease tau phosphorylation, which is expected to enhance MT assembly and axon elongation. Thus, overexpressing aPKC might rescue the polarity defects caused by ectopic MARK2 expression. However, ectopic aPKC was unable to rescue the defects caused by MARK2 (Fig. 4A and E). Surprisingly, coexpression of all three components of the PAR-3/PAR-6/aPKC complex did exhibit a rescue effect (Fig. 4A–C, and E). Approximately half of the cells expressing EGFP-MARK2 together with PAR-3/PAR-6/aPKC exhibited normal polarity, with at least one long process staining strongly for Tau1 (Fig. 4C and E). In contrast, only 10% of cells cotransfected with MARK2...
with the PAR-3 caused by the T595A mutant could not be rescued by cotransfection

In this study, we investigated the role of MARK in establishing neuronal polarity. We report that: (i) expression of a wild-type MARK2 in hippocampal neurons results in an increase of tau phosphorylation at S262 and a loss of neuronal polarity; these MARK2-induced phenotypes were rescued by coexpression of the PAR-3/PAR-6/aPKC polarity complex; (ii) reducing MARK2 levels in hippocampal neurons promotes axon outgrowth resulting in neurons containing several axons, rather than a single one, and reducing MARK2 levels rescues polarity defects induced by expression of a dominant-negative aPKC mutant; and (iii) perturbing endogenous MARK2 interactions with aPKC by overexpressing an MARK2 mutant (MARK2MC) abrogates neuronal polarity. Taken together, these results suggest that MARK2 inhibition by aPKC plays an active role in regulating neuronal polarity and, in particular, in regulating axon development. We propose that aPKC, when complexed with PAR-3 and PAR-6, negatively regulates MARK(s), which in turn leads to dephosphorylation of MAPs such as tau, and finally promotes the assembly of stable MTs and axon elongation.

Modulation of MT stability by phosphorylation of MAPs appears to be a major determinant for axon specification (3, 8, 11). MARK2 phosphorylates tau at S262 (23, 32), which reduces tau binding to MTs (33). Similar to the PAR-3/PAR-6/aPKC complex (15, 16, 34), both MARK2 and an inhibited form of MARK2, p-MARK2 (T595), distributed mainly in axons of hippocampal neurons at stage 3. In contrast, p-tau (S262) was concentrated in dendrites but absent from axons (Fig. 6) (28), indicating that MARK2 is inactivated in the developing axons. The findings that PAR-3/PAR-6/aPKC complex and aPKC activity are essential for the establishment of neuronal polarity (15, 16, 34) and that aPKC interacts with and negatively regulates MARK2 (26) led us to postulate that PAR-3/PAR-6/aPKC might function by negatively regulating MARK2 in neuronal polarization. The ability of aPKC to inhibit MARK2-mediated tau phosphorylation was demonstrated in HEK-293 cells (Fig. 2), PC12 cells (see Fig. 8), and primary neurons (Fig. 4).

Knockdown MARK2 levels with siRNA decreased tau phosphorylation at S262, induced formation of multiple axon-like neurites, and promoted axon outgrowth. These results imply that some fraction of MARK2, such as that in dendrites or soma, might be in an active state in which inhibition is sufficient to promote MT assembly and axon outgrowth.

Expression of an MARK2 mutant (MARK2MC) lacking a kinase domain but retaining aPKC-binding domain caused tau phosphorylation and polarity defects (Fig. 3), presumably by preventing aPKC from negatively regulating endogenous MARK2. These findings further support the conclusion that aPKC negatively regulates MARK2 to promote neuronal polarity. Additional support for this conclusion comes from the following observations: inhibition of aPKC in hippocampal neurons led to decreased levels of p-MARK2 (T595), an “inhibited” form of MARK2 (Fig. 2B); expression of the PAR-3/PAR-6/aPKC complex partially rescued the phenotype induced by overexpression of wild-type MARK2 but not the T595A phosphorylation mutant; and lastly, MARK2 knockdown partially rescued the polarity defects caused by expression of a dominant-negative aPKC mutant (Fig. 5).

Inhibition of GSK3β by the phosphatidylinositol 3-kinase/AKT pathway is essential for neuronal polarization (12, 13). The substrates of GSK3β include several MAPs, such as CRMP-2, MAP1B, tau, and Apc, in which phosphorylation prevents interaction with MTs (12, 35–38). Interestingly, GSK3β phosphorylation of tau requires prior phosphorylation by PAR-1 in Drosophila (32). It will be interesting to investigate whether GSK3β phosphorylation of tau also requires MARKs in the mammalian system. A recent study suggests a causal relationship between PAR-1 and GSK3β, whereby aPKCβ directly phosphorylates and activates MARK2, PAR-1 (39). Another possibility is that phosphatidylinositol 3-kinase/AKT/GSK3β pathway acts in parallel with the PAR-3/PAR-6/aPKC/MARK2 pathway in establishing neuronal polarity. Activation of either pathway may be sufficient to promote MT assembly and axon elongation.

Expression of PAR-1/MARK2 in neuroblastoma N2a cells promotes neurite outgrowth, whereas the formation of neurites is blocked if MARK2 is inactivated (23). In addition, PAR-1/MARK...
kinase was shown to be a positive regulator of MT-dependent transportation in the axons of RGC neurons (40) but a negative regulator of the cortical actin network via association with PAK5 (41). Local destabilization of F-actin may be a trigger for axonal specification (10). Interestingly, during epithelial polarization, reduced MARK2 levels are associated with weaker peripheral F-actin staining (20), raising an intriguing possibility that MARK2-siRNA-induced formation of multiple axons might be caused by local actin dynamics. The mechanisms that coordinate MT and actin cytoskeleton assembly and dynamics remain to be clarified.

Methods

Reagents, Antibodies, and Constructs. The antibodies used for immunostaining or immunoblotting were from Chemicon (MAP2, Tau1, and synapsin), Santa Cruz Biotechnology (PKCζ), BioSource International (p-tau (S262)), Calbiochem (tau), and Sigma (β-tubulin III). Rabbit anti-MARK2 antibody was generated by immunization with GST-hMARK2 (amino acids 531–670) as an antigen. Rabbit anti-p-MARK2 at T595 was reported in ref. 26. Double-stranded oligonucleotides targeting against MARK2 were synthesized by Shanghai GenePharma (Shanghai, China). The siRNA sequence was designed against a region starting from nucleotide 1180 or 1677 of rat MARK2 cDNA. The sequence from nucleotide 1677 is 5'-GAATGAACCTGGAAGCAGAA-3' (forward) and 5'-TTTGCTTCCAGTTCTC-3' (reverse). The sequence from nucleotide 1180 is 5'-CAGATGAAAACGCA-GAAGA-3' (forward) and 5'-TTTCGCGGTGTCTCCTG-3' (reverse). SAD plasmids were kindly provided by J. R. Sanes (Harvard University, Boston). EGFP-tag plasmids were gifts from K. S. Kosik (Harvard Medical School, Boston). PAR-3 and PAR-6 were gifts from I. G. Macara (University of Virginia, Charlottesville) (30). Flag-PKζ and PAR-1b/MARK2 (wild type and T595A mutant) constructs were introduced in ref. 26. MARK2 fragments were subcloned into BamHI and KpnI sites of pEGFP-N1. aPKC-N (amino acids 1–109) was subcloned into BamHI and EcoRI sites of the pKH3 vector.

Neuron Culture and Electroporation. Hippocampi of embryonic day-17 rat embryos were digested with 0.125% trypsin/EDTA for 20 min at 37°C, followed by trituration with pipettes in the plating medium (DMEM with 10% FBS). Dissociated neurons were transfected by using a nucleofector device (Ammax, Gaithersburg, MD). In some experiments, plating media were cotransfected with pEGFP with a ratio of 3:1. Control and transfected neurons were plated onto coverslips coated with poly-l-lysine (0.1 mg/ml) and laminin (0.05 mg/ml). After culturing for 4 h, media were changed to neuronal culture medium (neurobasal media containing 1% glutamate and 2% B27).

Immunohistochemistry. Neurons were washed with PBS, fixed in 4% paraformaldehyde at 4°C for 45 min, and incubated with 0.1% Triton X-100 in PBS for 10 min. After blocking with 10% goat serum in PBS at room temperature for 1 h or at 4°C overnight, neurons were incubated in primary antibodies at 4°C for 12 h and subsequently with Alexa Fluor 350-, FITC-, or rhodamine-conjugated secondary antibodies. Coverslips were mounted and examined by using a NeuroLucida system (Nikon).

Biochemical Characterization. Rat brains were homogenized in cold lysis buffer containing 50 mM Tris-HCl, pH 7.5, 150 mM NaCl, 1% Nonidet P-40, 0.5% sodium deoxycholate, and protease inhibitors. After clarification by centrifugation (100,000 × g; 1 h at 4°C), lysates were immunoblotted with indicated antibodies. HEK-293 cells were cultured in DMEM supplemented with 10% FBS and transfected with the standard calcium phosphate method. Cell lysates were prepared in the cold lysis buffer. Immunoprecipitation and immunoblotting were performed as described in ref. 42.

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