Region-Specific Phosphorylation of Rabphilin in Mossy Fiber Nerve Terminals of the Hippocampus

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In mossy fiber synapses of the CA3 region of the hippocampus, long-term potentiation (LTP) is induced presynaptically by activation of CAMP-dependent protein kinase A (PKA). Rab3A is a synaptic vesicle protein that regulates vesicle fusion and is essential for mossy fiber LTP. Rab3A probably acts via two effector proteins, rabphilin and RIM, of which rabphilin is an in vitro substrate for PKA. To test if rabphilin is phosphorylated in nerve terminals and if its PKA-dependent phosphorylation correlates with the PKA-dependent induction of LTP in mossy fiber terminals, we have studied the phosphorylation of rabphilin in synaptosomes isolated from the CA1 and CA3 regions of the hippocampus. Rabphilin was phosphorylated in both CA1 and CA3 synaptosomes. However, when we treated the CA1 and CA3 synaptosomes with forskolin (an agent that enhances PKA activity) or induced Ca\(^{2+}\) influx into synaptosomes with high K\(^+\), rabphilin phosphorylation was increased selectively in mossy fiber CA3 synaptosomes, but not in CA1 synaptosomes. In contrast, the phosphorylation of synapsin, studied as a control for the specificity of the region-specific phosphorylation of rabphilin, was augmented similarly by both treatments in CA1 and CA3 synaptosomes. These results reveal that the phosphorylation states of two synaptic substrates for PKA and CaM KII, rabphilin and synapsin, are regulated differentially in a region-specific manner, an unexpected finding because rabphilin and synapsin are similarly present in CA1 and CA3 synaptosomes and are colocalized on the same synaptic vesicles. The region-specific phosphorylation of rabphilin agrees well with the restricted induction of LTP by presynaptic PKA activation in mossy fiber, but not CA1, nerve terminals.

Key words: rab3; rabphilin; exocytosis; neurotransmitter release; synaptic transmission; synaptic vesicles; long-term potentiation; protein kinase A; CaM kinase II

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preparations used to study presynaptic function electrophysiologically are inaccessible biochemically. A second problem is that it is unclear which synaptic proteins are phosphorylated in nerve terminals in correlation with functional changes. This gap in understanding is most striking for synapsins and rabphlin, which are the only currently known presynaptic PKA and CaM KII substrates, but for which the phosphorylation in nerve terminals either has not been studied yet (rabphilin) or has not been associated directly with a synaptic plasticity (synapsins). In the current study, we have investigated the phosphorylation of rabphilin and synapsins in synaptosomes from hippocampal synapses with distinct properties, regular synapses from the CA1 region and large mossy fiber synapses from the CA3 region. Our results demonstrate that rabphilin is phosphorylated in a stimulation-dependent and region-specific manner that correlates with mossy fiber LTP. This suggests a mechanism of presynaptic regulation that uses region-specific changes in the phosphorylation states of generally distributed synaptic proteins.

MATERIALS AND METHODS

Preparation of synaptosomes. CA1 and CA3 regions were dissected from rat hippocampus, using the minor and the hippocampal fissures as landmarks. Large mossy fiber synaptosomes were prepared by homogenizing the CA3 region manually in 1 mM MgSO4, 0.3 M sucrose, and 15 mM HEPES-NaOH, pH 7.4, to preserve the complex structure and large size of the mossy fiber terminals (Hajos et al., 1975; Terrian et al., 1988). The homogenate was passed through a series of nylon filters (100, 60, and 40 μm mesh size) and centrifuged at 900 × g for 10 min to selectively pellet large synaptosomes. The pellet was resuspended in 18% (w/v) Ficol and 0.3 M sucrose and centrifuged at 16,000 × g for 20 min to separate the synaptosomes that remained in suspension from contaminating nuclei. Synaptosomes were diluted with 4 vol of 0.3 M sucrose and sedimented by centrifugation at 15,000 × g for 20 min to remove the Ficol before use. Small synaptosomes were prepared from the CA1 region by gradient centrifugation by a method modified from Nagy and Delgado-Escueta (1984). The crude mitochondrial fraction (P2) was resuspended in 8.5% (w/v) Percoll suspended in 0.25 M sucrose and 5 mM HEPES-NaOH, pH 7.4, and layered on top of an 12%/20% Percoll step gradient in the same buffer. After centrifugation at 18,000 × g for 30 min, synaptosomes were recovered from the 12%/20% Percoll interface. Percoll was removed by the addition of 30 vol of 0.32 M sucrose and centrifugation at 18,000 × g for 20 min. In a typical experiment 2 mg of CA1 and 4 mg of CA3 synaptosomal protein were isolated from seven rat brains.

Phosphorylation measurements. Synaptosomes (1 mg protein) were incubated with 32P-orthophosphate (1 mCi) for 1 hr at 37°C in phosphate-free aerated (95% O2/5% CO2) Krebs–Henseleit–HEPES buffer; composition (in mM): NaCl 116, KCl 3.5, CaCl2 1.25, MgSO4 1.2, KH2PO4 1.2, NaHCO3 25, HEPES-NaOH 5 at pH 7.4, and glucose 11.5) containing 0.1 mM CaCl2. The tubes were flushed with 95% O2/5% CO2 every 15 min. Free 32P-orthophosphate mark serum (1% v/v final concentration) was added. Rabphilin was immunoprecipitated with antibodies against the recombinant GST-rabphilin fusion protein (I734; Li et al., 1994), and synapsins were raised with an antibody against the N-terminal sequence of synapsins (E028; Rosahl et al., 1995). Antigen–antibody complexes were recovered by incubation with 50 μl of a 50% slurry of protein A-Sepharose beads for 45 min and subsequent centrifugation at 1000 × g for 5 min. The beads were washed four times in 10 mM HEPES-NaOH, pH 7.4, 0.15 M NaCl, 1 mM EDTA, and 0.5% Triton X-100; resuspended in SDS sample buffer; and analyzed by SDS-PAGE. After staining and destaining, phosphoproteins were visualized by autoradiography. 32P incorporation was quantified with a PhosphorImager (Molecular Dynamics, Sunnyvale, CA). Data from multiple experiments performed in duplicate were pooled, and the significance of changes was evaluated statistically with the Kruskal–Wallis one-way ANOVA, followed by Dunnett’s test for pairwise multiple comparisons, with a significance level of p < 0.05.

RESULTS

Purification of CA1 and CA3 synaptosomes from the hippocampus

To study the differential regulation of hippocampal synapses biochemically, we purified synaptosomes from the CA1 and CA3 regions of the hippocampus. We dissected the CA1 and CA3 regions from rat brains, isolating regular “small” synaptosomes from the CA1 region by standard techniques and large mossy fiber synaptosomes from the CA3 region by making use of their large size (see Materials and Methods). To determine whether the protein composition of the two types of synaptosomes is similar, we analyzed them by immunoblotting with antibodies to several presynaptic proteins (Fig. 1). No major differences in the relative levels of different proteins were found.
Protein phosphorylation in CA1 and CA3 synaptosomes

To gain insight into potential differences between CA1 and CA3 synaptosomes in the phosphorylation of proteins by PKA, we first analyzed the overall pattern of protein phosphorylation in 32P-labeled synaptosomes. Synaptosomes were analyzed by SDS-PAGE and autoradiography either under control conditions or after treatment with forskolin, elevated K\(^+\), or a combination of forskolin and elevated K\(^+\). Forskolin was used because it activates adenylate cyclase, thereby increasing cAMP levels and PKA activity. Elevated K\(^+\) was used because it causes Ca\(^{2+}\) influx into synaptosomes, thereby activating CaM K11 and other kinases. Thus the two treatments activate two different but overlapping signal transduction pathways in synaptosomes, one of which (the PKA pathway) is required for CA3 LTP, but not CA1 LTP.

No major differences in the phosphoprotein pattern of CA1 and CA3 synaptosomes were observed under resting conditions, providing additional evidence that they are similar in composition (Fig. 2). Forskolin did not induce major alterations in the phosphorylation patterns of either CA1 or CA3 synaptosomal proteins. K\(^+\) elevation increased phosphorylation of a protein in the 10 kDa molecular weight range in both types of synaptosomes but otherwise caused only minor changes (arrow in Fig. 2). We also analyzed synaptosomal protein phosphorylation by two-dimensional gel electrophoresis after various stimulation conditions to confirm these findings (Garrels, 1979). Although the two-dimensional gels improved resolution, we again observed no differences between CA1 and CA3 synaptosomes (data not shown). These results suggest that CA1 and CA3 synaptosomes are similar in composition and that forskolin treatment and K\(^+\) depolarization do not cause major changes in the phosphorylation of abundant proteins.

Phosphorylation of rabphilin and synapsins in CA1 and CA3 synaptosomes

We next studied whether rabphilin is phosphorylated in the two types of synaptosomes as a system resembling in vivo conditions. For these and the following experiments, we used synapsins as well characterized synaptic phosphoproteins as positive controls. We immunoprecipitated rabphilin and synapsins from resting 32P-labeled synaptosomes and from synaptosomes that were treated with okadaic acid. Okadaic acid was applied at a concentration of 50 μM at which it is an inhibitor of serine/threonine protein phosphatases 1, 2A, and 2B. At this concentration okadaic acid nonspecifically increases protein phosphorylation. The experiments with okadaic acid revealed that rabphilin and synapsins were phosphorylated in both CA1 and CA3 synaptosomes and that okadaic acid greatly increased the phosphorylation states of both proteins in the two types of synaptosomes (Fig. 3). Thus rabphilin is phosphorylated in synaptosomes similar to synapsins. However, phosphorylation and immunoprecipitation experiments can be quite variable because they involve multiple procedures, and single experiments are insufficient for evaluating magnitudes of changes. Therefore, we quantitated the degree of rabphilin and synapsin phosphorylation in multiple independent experiments with a phosphoimager. The increase in phosphorylation induced by okadaic acid was calculated for each experiment relative to baseline phosphorylation.

Using this analysis, we found that okadaic acid increased both rabphilin and synapsin phosphorylation three- to fourfold in both types of synaptosomes (Fig. 4). These data quantitatively confirm the qualitative impression from the experiment in Figure 3, namely that rabphilin and synapsins are phosphoproteins in synaptosomes, the phosphorylation of which constantly turns over and therefore is enhanced by the addition of a phosphatase inhibitor. The overall phosphorylation signal is higher for synapsins than for rabphilin, presumably because synapsins are more abundant. However, the increase in phosphorylation induced by okadaic acid is similar for rabphilin and synapsins and not dramatically different between CA1 and CA3 synaptosomes. This indicates that at resting conditions the turnover rate of phosphorylation is comparable for the two proteins and the two types of synaptosomes.
Forskolin and high K\(^+\) enhance rabphilin phosphorylation in CA3, but not in CA1, synaptosomes

Next we investigated the regulation of rabphilin phosphorylation by PKA (activated by forskolin treatment) and by Ca\(^{2+}\)-dependent protein kinases (activated by Ca\(^{2+}\) influx triggered by membrane depolarization in solutions with 20 mM K\(^+\)). \(^{32}\)P-labeled synaptosomes from the CA1 and CA3 regions were treated with four conditions: (1) control, (2) forskolin, (3) K\(^+\) depolarization, and (4) forskolin in combination with K\(^+\) depolarization. We then immunoprecipitated rabphilin from the treated synaptosomes and quantified its phosphorylation by SDS-PAGE and phosphoimager detection. To ensure that the immunoprecipitations were equally effective under various treatment conditions, we confirmed the presence of rabphilin in the immunoprecipitates by immunoblotting (data not shown). Because these experiments involved multiple procedures that might fluctuate randomly between experiments, we quantified phosphorylations in multiple independent determinations. In each experiment the phosphorylation of rabphilin was measured relative to baseline phosphorylation, and data from multiple experiments were analyzed statistically.

Forskolin induced a dramatic increase in rabphilin phosphorylation in CA3 synaptosomes (260–400% increase), suggesting that it is a physiological substrate for PKA (activated by forskolin treatment) and by Ca\(^{2+}\) (data not shown). The combination of forskolin with K\(^+\) depolarization caused an additive increase in phosphorylation in the CA3 synaptosomes but only a small effect in CA1 synaptosomes. These observations suggest that rabphilin is a substrate for PKA and CaM KII not only in vitro but also in synaptosomes, a system resembling nerve terminals of living neurons.

Strikingly, forskolin had no significant effect on rabphilin phosphorylation in CA1 synaptosomes. The stimulatory effect of K\(^+\) on rabphilin phosphorylation also was restricted to synaptosomes from the CA3 region, and no effect was observed in CA1 region synaptosomes (Fig. 5). These results reveal that the phosphorylation of rabphilin is regulated differentially in different types of nerve terminals (CA1 vs CA3).

Forskolin and depolarization-induced Ca\(^{2+}\) influx stimulate phosphorylation of synapsins

The fact that forskolin and Ca\(^{2+}\) influx increase rabophilin phosphorylation in a region-specific manner is striking because okadaic acid increased phosphorylation of rabphilin in both types of synaptosomes (Fig. 5). This raises the question of whether any unknown reason forskolin and Ca\(^{2+}\) influx have been unable to activate kinases in the CA1 synaptosomes. To evaluate this concern, we analyzed the effect of forskolin and Ca\(^{2+}\) influx on the phosphorylation state of synapsins in both types of synaptosomes (Fig. 6). Our results show that both forskolin and K\(^+\) elevation stimulated synapsin phosphorylation and stimulated it equally in CA1 and CA3 synaptosomes. The increase in synapsin phosphorylation was relatively small (35–50%), as compared with the...
increase in rabphilin phosphorylation observed in CA3 synaptosomes (260–400%). However, the increase in synapsin phosphorylation under these mild stimulation conditions is statistically significant ($p < 0.05$) and similar to that observed by others previously (Wang et al., 1988). The fact that synapsin phosphorylation is increased equally in CA1 and CA3 synaptosomes demonstrates that the CA1 synaptosomes are responsive to stimulation and that the lack of an increase in rabphilin phosphorylation under our treatment conditions is not attributable to a failure of these synaptosomes to respond.

**DISCUSSION**

In presynaptic mossy fiber terminals of the CA3 region of the hippocampus, PKA activation induces a presynaptic form of LTP, which we refer to as CA3 LTP (Huang et al., 1994; Weissskopf et al., 1994). Recent studies have shown that, in mice lacking the synaptic vesicle protein rab3A, CA3 LTP cannot be induced normally (Castillo et al., 1997). This result directly proves the participation of the molecular machinery for vesicular exocytosis in CA3 LTP. It suggests that rab3A is, in an unknown manner, required for PKA-dependent long-term potentiation of neurotransmitter release. Rab3A itself is not a PKA substrate, suggesting that PKA does not act directly on rab3A. However, rab3A is thought to act by binding to effector proteins in a GTP-dependent manner, and one of its two known effectors, rabphilin, was shown previously to be a substrate for PKA in vitro (Fykse et al., 1995).

In the present study we set out to address two questions: (1) Is rabphilin also phosphorylated by PKA in a system resembling nerve terminals of living neurons? (2) Is there a difference in the phosphorylation of rabphilin between CA1 and CA3 nerve terminals that correlates with the ability of PKA to induce LTP in mossy fiber synapses of the CA3 region, but not in Schaffer collateral/commissural fiber synapses of the CA1 region?

To resolve these questions, we have isolated synaptosomes, pinched-off nerve endings, from the CA1 and CA3 regions of the hippocampus and have analyzed the phosphorylation of rabphilin in these synaptosomes. Synapsin phosphorylation was studied as a control. We quantitatively compared the phosphorylation of rabphilin and synapsins to baseline phosphorylation states under four treatment conditions: (1) okadaic acid as a relatively nonspecific phosphatase inhibitor to induce a general increase in phosphorylation states, (2) forskolin as an activator of adenyl cyclase to increase cAMP levels and PKA activity, (3) elevated $K^+$ to trigger $Ca^{2+}$ influx into the synaptosomes to activate $Ca^{2+}$-dependent kinases, and (4) a combination of forskolin and $K^+$ treatment.

Our data result in two basic conclusions. (1) Rabphilin is phosphorylated in synaptosomes, and its phosphorylation is increased several-fold by activations of PKA and by $Ca^{2+}$ influx. This suggests that it constitutes a physiological substrate for PKA and for $Ca^{2+}$-activated kinases, probably CaM Ki. (2) Although rabphilin is phosphorylated and its phosphorylation is increased by a generic phosphatase inhibitor in both CA1 and CA3 synaptosomes, forskolin and $Ca^{2+}$ influx enhance rabphilin phosphorylation only in CA3, but not in CA1, synaptosomes. The region-specific regulation of rabphilin phosphorylation parallels that of the action of PKA in inducing CA3 LTP. It is unlikely to be artificial, because synapsin phosphorylation was increased similarly in both types of synaptosomes with forskolin and $K^+$ stimulation.

The region-specific phosphorylation of rabphilin is surprising because phosphorylated rabphilin and active PKA are present in both types of synaptosomes. These results reveal an unexpected regional specialization of protein phosphorylation in nerve terminals, the first demonstration of region-specific phosphorylation of a synaptic protein. The region-specific phosphorylation of rabphilin is particularly striking in view of the behavior of synapsins, which are colocalized with rabphilin on the same vesicles and are substrates for the same kinases but are still not subject to the same region-specific phosphorylation.
Two questions arise: (1) What is the mechanism that allows the region-specific phosphorylation of rabphilin? (2) What does this phosphorylation mean for CA3 LTP? Although we have no definitive answer to either question at present, the fact that forskolin and K⁺ depolarization have similar effects on synapsin phosphorylation in CA1 and CA3 synaptosomes suggests that PKA and CaMK kinases are similarly present in both types of nerve terminals. Thus the selective phosphorylation of rabphilin in CA3 synaptosomes could be attributable to a compartmentalization of the kinases, a possibility raised by observations with other phosphorylation events (for review, see Faux and Scott, 1996). Alternatively, it could be attributable to differential activities of phosphatases in the two types of nerve terminals. Independent of which mechanism is operative, the net result is a differential phosphorylation state of rabphilin in the two types of nerve terminals, which thereby adds a layer of regulation on top of activation of protein phosphorylation and protein expression.

As regarding the question of the functional significance of the differential phosphorylation of rabphilin in CA3 versus CA1 synaptosomes, the correlation of this phenomenon with the selective induction of PKA-dependent LTP in mossy fiber nerve terminals (CA3 LTP) and the requirement for rab3A in the induction is intriguing. This suggests the possibility that rab3A may be required for CA3 LTP because it interacts with rabphilin and reversibly recruits rabphilin to nerve terminals (Li et al., 1994; Stahl et al., 1996). At present this possibility cannot be readily tested because forskolin has additional actions besides activating PKA and inducing LTP in mossy fiber nerve terminals (Hosi et al., 1988; Huang et al., 1995; Castillo et al., 1997). These parallel actions are unimpaired in the rab3A knock-out mice so that, although CA3 LTP can no longer be induced, forskolin still enhances neurotransmitter release. Therefore so that rabphilin phosphorylation can be correlated with LTP, studies on rabphilin knock-outs will be required. Nevertheless, the specificity for the CA3 region of the PKA-dependent phosphorylation of rabphilin and of the PKA-dependent induction of LTP is suggestive of a role of rabphilin phosphorylation in CA3 LTP. In addition, the region-specific phosphorylation of rabphilin implies that the phosphorylation states of substrate proteins can be regulated locally if the kinases and substrates are present ubiquitously, even if the substrates are colocalized on the same organelle.

REFERENCES


