Targeting Protein Phosphatase 1 (PP1) to the Actin Cytoskeleton: the Neurabin I/PP1 Complex Regulates Cell Morphology

Carey J. Oliver,1 Ryan T. Terry-Lorenzo,1 Elizabeth Elliott,2 Wendy A. Christensen Bloomer,1 Shi Li,1 David L. Brautigan,2 Roger J. Colbran,3 and Shirish Shenolikar1*

Department of Pharmacology and Cancer Biology, Duke University Medical Center, Durham, North Carolina 27710;1 Center for Cell Signaling, University of Virginia School of Medicine, Charlottesville, Virginia 22908;2 and Department of Molecular Physiology and Biophysics, Vanderbilt University School of Medicine, Nashville, Tennessee 37232

Received 23 January 2002/Returned for modification 6 March 2002/Accepted 25 March 2002

Neurabin I, a neuronal actin-binding protein, binds protein phosphatase 1 (PP1) and p70 ribosomal S6 protein kinase (p70S6K), both proteins implicated in cytoskeletal dynamics. We expressed wild-type and mutant neurabins fused to green fluorescent protein in Cos7, HEK293, and hippocampal neurons. Biochemical and cellular studies showed that an N-terminal F-actin-binding domain dictated neurabin I localization at actin cytoskeleton and promoted disassembly of stress fibers. Deletion of the C-terminal coiled-coil and sterile alpha motif domains abolished neurabin I dimerization and induced filopodium extension. Immune complex assays showed that neurabin I recruited an active PP1 via a PP1-docking sequence. Mutation of the PP1-binding motif or PP1 inhibition by okadaic acid and calyculin A abolished filopodia and restored stress fibers in cells expressing neurabin I. In vitro and in vivo studies suggested that the actin-binding domain attenuated protein kinase A (PKA) phosphorylation of neurabin I. Modification of a major PKA site, serine-461, impaired PP1 binding. Finally, p70S6K was excluded from neurabin I/PP1 complexes and required the displacement of PP1 for recruitment to neurabin I. These studies provided new insights into the assembly and regulation of a neurabin I/PP1 complex that controls actin rearrangement to promote spine development in mammalian neurons.

Cross talk between protein kinases and phosphatases regulates synaptic strength and information processing in mammalian brain (33). Prior studies identified protein phosphatase 1 (PP1) as a key regulator of activity-dependent changes in synaptic function underlying the two major forms of plasticity known as long-term potentiation (3) and long-term depression (LTD) extensively studied for hippocampal neurons (19). LTD-inducing stimuli promoted distribution of PP1 to dendritic spines (18), where it associated with the actin-rich cytoskeleton, consistent with their ability to bind polymerized F-actin. More recent studies showed that the cytoskeletal assembly of neurabins was highly dynamic and regulated by phosphorylation of a neurabin I/PP1 complex (20) and suggested that NrbI was a multifunctional protein scaffold that regulated both membrane and cytoskeletal functions.

By immunohistochemistry, NrbII was localized to dendritic spines and thus called spinophilin (1). In contrast, NrbI was present in both spines and growth cones (20). Subcellular fractionation showed that both neurabins are present in highly purified preparations of PSD (25, 32) and growth cones (R. T. Terry-Lorenzo and S. Shenolikar, unpublished observations). Ectopic expression of NrbI (20) and NrbII (25) in cultured cells demonstrated that both proteins localized to actin cytoskeleton, consistent with their ability to bind polymerized F-actin. More recent studies showed that the cytoskeletal association of neurabins was highly dynamic and regulated by growth factors (29) and small GTPases (25). This suggested that, in response to physiological signals, neurabins delivered signaling proteins, such as PP1 and p70S6K, to the actin cytoskeleton to control cell morphology. However, to date the role of NrbI signaling in cells has not been investigated.

Disruption of the mouse spinophilin/NrbII gene resulted in significant deficits in PP1 signaling required for LTD. Changes in neuronal morphology, specifically an overabundance of spines (9), were noted in the hippocampus of young NrbII-null mice. In contrast, antisense depletion of NrbI inhibited neurite outgrowth (20) and suggested distinct roles for the two neurabin isoforms in neurons. We have undertaken a combined molecular and cell biological analysis of rat NrbI to define its role in signal transduction and cell morphology. Our studies examined the contribution of specific structural deter-
Materials and Methods

Phosphorylation b was purchased from Calzyme. Actin Binding Kit BK001 was from Cytoskeleton. [γ-32P]ATP and [32P]orthophosphoric acid were purchased from NEN. Adenosine 3′,5′-cyclic monophosphorothioate, Sp isomer–triethylammonium salt (Rp-cAMPs), adenosine 3′,5′-cyclic monophosphorothioate, Rp isomer–triethylammonium salt (Rp-cAMPS), latrunculin B; okadaic acid and calycin A were from Calbiochem. Anti-p70S6K antibody was obtained from Santa Cruz Biotechnology, anti-green fluorescent protein (anti-GFP) was from Research Diagnostics, and anti-PP1 was from Transduction Laboratories.

Expression of Nrb in Escherichia coli. His-Nrb (1-615), His-Nrb (103-615), His-Nrb (615-1095), and His-Nrb (1095-1095) were generated by PCR with full-length rat Nrb cDNA as template and cloned into pRSET-B (Invitrogen). BL2(DE3)Lyss competent cells (Stratagene) were transformed with plasmid pETDuet-1 (Novagen). Recombinant E.coli BL21(DE3)pLysS competent cells (Stratagene) were transformed with plasmid pETDuet-1, which was purchased from Novagen (Madison, WI). HeLa and COS7 cells were maintained in Dulbecco modified Eagle medium (DMEM) supplemented with 10% fetal bovine serum and 100 U/ml of penicillin, 100 μg/ml of streptomycin, 2 mM of L-glutamine, and 1% antibiotic-antimycotic mixture at 37°C. Images were captured every 150 s for 1 h with a Hamamatsu Orca II cooled charge-coupled device camera controlled by live software (Inovision).

Immune complex phosphatase assays. GFP-Nrb immunoprecipitates adsorbed to protein G-Sepharose were washed in NETN buffer as described above followed by 50 mM Tris-HCl (pH 7.5)–1 mM EDTA–0.1% (vol/vol) β-mercaptoethanol and assayed for protein phosphatase activity with [32P]labeled phospho-tyrosine at 37°C for 30 min (26).

Metabolic labeling of HEK293 cells. HEK293 cells expressing GFP-Nrb were incubated in DMEM lacking sodium phosphate and sodium pyruvate but containing 12.5 μCi of [32P]orthophosphoric acid for 2 h. Cells werestimulated with 100 μM Sp-cAMPS or Rp-cAMPS for 15 min, washed twice with ice-cold PBS, and lysed in 500 μl of RIPA buffer containing 50 mM sodium fluoride for 15 min on ice. Lysates were clarified by centrifugation at 16,000 × g for 5 min at 4°C. Supernatants were precleared with protein G-Sepharose (1 h at 4°C), which was removed by centrifugation at 100,000 × g for 1 h at 4°C. The supernatants were incubated with anti-GFP for 1 h at 4°C followed by protein G-Sepharose (1 h). These beads were washed three times with NETN-250 (10 mM Tris-HCl [pH 8.0], 250 mM NaCl, 0.5% NP-40, 1 mM EDTA) and twice with NETN prior to the addition of SDS sample buffer and subjected to SDS–7.5% (wt/vol) PAGE and autoradiography.

F-Actin binding assays. F-actin binding was assayed with the Actin Binding Protein Biochem Kit BK001. Briefly, His-Nrb (7 μM) was incubated with F-actin, polymerized in a polymerization buffer, for 30 min at 24°C and centrifuged through a sucrose cushion at 150,000 × g for 1.5 h at 24°C. Supernatants and pellets were subjected to SDS–10% (wt/vol) PAGE, and the proteins were stained with Coomassie blue.

Pull-downs from brain extracts. Rat brain cortex (Pel-Freez) was homogenized with 25 volumes of 50 mM Tris-HCl (pH 7.5) containing 5 mM EDTA, 5 mM EGTA, 10 mM NaCl, 1% deoxycholate, 1 mM PMSF, 1 mM benzamidine, 1 μg of leupeptin/ml, and 1 μg of aprotinin/ml. The homogenate was centrifuged at 100,000 × g for 1 h, and the supernatant was dialyzed against 50 mM Tris-HCl (pH 7.5) containing 1 mM EDTA, 1 mM EGTA, 10 mM NaCl, 0.1 mM PMSF, and 0.1 mM benzamidine. Lysates (4 mg of total protein) were precleared with Ni−NTA agarose before they were incubated with 20 μg of His-tagged Nrb bound to Ni−NTA agarose for 1 h at 4°C. Beads were washed four times with PBS containing 5 mM imidazole, and bound proteins were eluted with 50 μl of SDS buffer for SDS–10% (wt/vol) PAGE and Western immunoblotting.

Pull-downs of brain extracts with microcystin-LR-Sepharose were performed as previously described (6).

In vitro phosphorylation of neurabin. His-Nrb proteins (10 μM) were incubated with protein kinase A (PKA) purified from bovine heart (2) in 50 μl of 10 mM Tris-HCl (pH 7.5) containing 300 μM dithiothreitol, 1 mM MgCl2, 100 μM ATP, and 0.03 μM of [γ-32P]ATP at 37°C. Aliquots (4 μl) were removed at intervals, and 50 μl of bovine serum albumin (10 mg/ml) and 200 μl of 20% trichloroacetic acid were added. Samples incubated at 4°C for 5 min were centrifuged at 13,000 × g for 5 min, and 32P incorporation into the trichloroacetic acid-precipitated fraction was determined by Cerenkov counting.

Results

N-terminal F-actin-binding domain localizes Nrb to actin cytoskeleton. An N-terminal actin-binding region was identified in rat neurabin (20,25) that was partly conserved in Xenopus laevis neurabin (accession no. AF363388) but absent from Caenorhabditis elegans (accession no. U80487) and Drosophila melanogaster neurabin (11). Thus, we analyzed hexahistidine-tagged recombinant rat Nrb polypeptides for in vitro F-actin binding (Fig. 1A). Nrb (1-615), like recombinant α-α-
tinin, a known F-actin-binding protein, was completely sedimented with polymerized F-actin. In contrast, NrbI (103-615) was largely soluble, with a very small amount of NrbI (103-615) consistently bound to F-actin. Earlier work showed that an N-terminal fragment (144 amino acids) of rat NrbI bound F-actin in vitro (20). A longer NrbI (1-210) polypeptide, however, bound F-actin more effectively. Moreover, myc-tagged NrbI (145-1095) expressed in Cos7 cells was partially localized at the actin cytoskeleton (20). This suggested that more extended sequences may dictate NrbI localization with the cytoskeleton.

We expressed GFP, GFP-NrbI (1-1095), and GFP-NrbI (286-1095). Cells were fixed with paraformaldehyde following transfection, permeabilized, and stained with rhodamine-phalloidin. Bar, 30 μm.

FIG. 1. N-terminal domain localizes GFP-NrbI to actin cytoskeleton. (A) Domain structure of rat NrbI. The protein interaction domains highlighted include an actin-binding domain (ABD), the PSD95/Dlg/ZO-1 domain (PDZ), the coiled-coil domain (CC), and the SAM domain. The PP1-binding domain (black box) is characterized by a canonical KIKF sequence. Also shown is the association of His-tagged rat NrbI (1-615) and rat NrbI (103-615) in vitro with polymerized F-actin. α-Actinin, a known F-actin-binding protein, was used as a control. Proteins were incubated in the presence (+) or absence (−) of F-actin in the actin polymerization buffer (cytoskeleton). Samples were centrifuged at 150,000 × g through a sucrose cushion, and particulate (P) and supernatant (S) fractions were analyzed by SDS–10% (wt/vol) PAGE and stained with Coomassie blue. Open arrowheads indicate the test proteins, and closed arrowheads indicate actin. (B) Fluorescence micrographs of Cos7 cells expressing GFP, GFP-NrbI (1-1095), or GFP-NrbI (286-1095). Cells were fixed with paraformaldehyde following transfection, permeabilized, and stained with rhodamine-phalloidin. Bar, 30 μm.
(286-1095) in Cos7 cells. GFP was widely distributed throughout the cell (Fig. 1B), and little overlap of GFP, seen as yellow, with F-actin-containing stress fibers or cortical actin cytoskeleton, stained by rhodamine-conjugated phalloidin, was seen. In contrast, full-length GFP-NrbI (1-1095) localized predominantly with actin-based structures, highlighted by the merged images. Interestingly, stress fibers in cells expressing GFP-NrbI (1-1095) were largely collapsed, as emphasized by actin staining in adjacent nontransfected cells. Treatment of cells with latrunculin B, an actin-depolymerizing agent, resulted in the redistribution of GFP-NrbI (1-1095) to diffuse intracellular sites, indicating that NrbI localization was dictated by the polymerized actin (data not shown). Similar results were obtained for HEK293 cells in which both GFP-NrbI (1-1095) and NrbI (1-1095)-GFP with GFP fused to the C terminus of NrbI induced a rounded morphology concomitant with the loss of stress fibers (data not shown). GFP-NrbI (286-1095), which lacked F-actin binding, was thus largely distributed to intracellular sites (Fig. 1B), and the cells retained their network of stress fibers. Neither GFP-NrbI (1-1095) nor GFP-NrbI (286-1095) entered the cell nucleus. Association with cytoskeleton was also confirmed by lysing cells in buffers containing 0.75% Triton X-100, which maintains an intact cytoskeleton (21), and centrifugation at 150,000 × g on a layer of 6% sucrose. Immunoblotting these fractions showed that GFP-NrbI (1-1095) bound exclusively to actin cytoskeleton, while both GFP and GFP-NrbI (286-1095) were soluble (data not shown). This established that F-actin binding was essential for both NrbI localization with the cytoskeleton and disassembly of stress fibers.

**C-terminal sequences in NrbI mediate dimerization.** Gel filtration of recombinant neurabin (20, 25) and immunoprecipitation of individual neurabin from extracts (16) suggested that NrbI formed homodimers and heterodimers with NrbII. However, the structural basis for neurabin dimerization which potentially contributes to its in vitro actin-bundling activity (20) remained undefined. Studies of other proteins containing coiled-coil (14) and SAM (27) domains suggested these domains as excellent candidates for NrbI dimerization. Thus, we undertook pull-downs from rat brain extracts with His-tagged NrbI as bait (Fig. 2A). To ensure direct association of the recombinant proteins with brain neurabins, we deleted the N-terminal 103 amino acids representing the actin-binding domain (Fig. 1A). Both His-NrbI (103-1095) and His-NrbI (615-1095) bound a 190-kDa polypeptide representing rat NrbI and, to a lesser extent, also bound NrbII. His-NrbI (1-615) failed to sediment either rat neurabin, providing the first direct evidence that C-terminal sequences in NrbI mediated homodimerization and heterodimerization.

To investigate the importance of NrbI dimerization, we expressed GFP-NrbI (1-552) and GFP-NrbI (553-1095) in Cos7 cells. As noted above, full-length GFP-NrbI (1-1095) localized with the actin cytoskeleton and resulted in collapse of stress fibers. GFP-NrbI (1-552) also localized with the actin cytoskeleton, and its distribution at the cell periphery was even more prominent (Fig. 2B). This was confirmed by biochemical fractionation as described above (data not shown). Cells expressing GFP-NrbI (1-552) not only demonstrated the loss of stress fibers but also projected extensive filopodia. GFP-NrbI (553-1095), representing the isolated dimerization domain, largely failed to localize with the cytoskeleton and was distributed in the cytoplasm. Cells expressing GFP-NrbI (553-1095) retained an intact stress fiber network.

GFP-NrbI (1-552) induced even more striking filopodia in HEK293 cells (Fig. 2C). Time-lapse video microscopy of living cells focused on the GFP-NrbI-containing filopodia seen in cells expressing GFP-NrbI (1-1095) and GFP-NrbI (1-552). Expression of wild-type (WT) GFP-NrbI (1-1095) resulted in the projection of short, highly dynamic filopodia that failed to extend further over the 10-min time frame. In contrast, not only did GFP-NrbI (1-552)-induced filopodia continue to extend over this time period (arrows in Fig. 2C) but also many of these projections remained stable for 60 min. These studies suggested that the dynamics of NrbI dimerization dictated its effects on cell morphology, promoting dissolution of stress fibers and filopodium extension.

**PP1 recruitment required for NrbI-induced filopodia and stress fiber disassembly.** GFP, GFP-NrbI (1-1095), and GFP-NrbI (1-552) expression in HEK293 cells had no effect on overall PP1 levels (Fig. 3A). Immunoprecipitation with anti-GFP monoclonal antibody showed that, despite its much higher expression, GFP alone bound only a trace of PP1 (Fig. 3A). In contrast, immunoprecipitates containing GFP-NrbI (1-1095) and GFP-NrbI (1-552) contained readily detectable PP1. Quantitative analysis of the immunoblots suggested that GFP-NrbI (1-552) bound PP1 more effectively than did WT GFP-NrbI (1-1095) (data not shown). Substituting Alanines for 457KIKF460, a proposed PP1-binding motif, eliminated PP1 association with GFP-NrbI (1-552, 4A).

Anti-GFP immunoprecipitates were also assayed for PP1 activity with the standard substrate, phosphorylase a. Extensive washes of anti-GFP immunoprecipitates with either NETN or assay buffer failed to release PP1 (data not shown), and the immune complex assays established for the first time that the NrbI-bound PP1 is an active protein phosphatase. When corrected for equivalent amounts of protein, immunoprecipitates containing GFP-NrbI (1-552) consistently contained 50 to 60% more phosphatase activity than did GFP-NrbI (1-1095) (Fig. 3B). As anticipated, immunoprecipitates containing GFP-NrbI (1-552, 4A) possessed little phosphatase activity, similar to GFP alone. These data showed that PP1 utilized the 457KIKF460 sequence for NrbI binding and bound GFP-NrbI (1-552) more effectively than GFP-NrbI (1-1095).

To confirm the identity of the NrbI-bound phosphatase, immune complex assays were performed with GFP-NrbI (1-552) in the presence of increasing concentrations of okadaic acid (Fig. 3C). Okadaic acid (5 nM), sufficient to inhibit most PP2A-like phosphatases, had no effect on GFP-NrbI (1-552)-bound phosphatase, and concentrations exceeding 100 nM, up to 1 μM, were needed to fully inhibit the enzyme, consistent with its being exclusively PP1 (5).

As indicated above, GFP-NrbI (1-552) expression collapsed stress fibers and induced filopodia in Cos7 cells (Fig. 3D). In contrast, the non-PP1-binding GFP-NrbI (1-552, 4A), while still localized to the actin cytoskeleton, failed to induce filopodia, and stress fibers remained intact in these cells. Similarly, in HEK293 cells, GFP-NrbI (1-552) elicited striking filopodia (Fig. 3E), which were completely eliminated in GFP-NrbI (1-552, 4A)-expressing cells, which possessed a rounded morphology similar to cells expressing WT GFP-NrbI (1-1095) (data...
FIG. 2. C-terminal sequences mediate dimerization of Nrbl. The Nrbl structure is schematically shown as in Fig. 1. (A) Proteins sedimented from rat brain deoxycholate extracts by 20 μg of His-Nrbl (103-1095), His-Nrbl (103-615), and His-Nrbl (615-1095) bound to Ni\(^{2+}\)-NTA agarose were subjected to SDS–10% (wt/vol) PAGE and immunoblotted with anti-Nrbl and anti-NrblII antibodies. (B) Fluorescence micrographs of Cos7 cells expressing GFP-Nrbl (1-1095), GFP-Nrbl (1-552), or GFP-Nrbl (553-1095). Cells were fixed with paraformaldehyde following transfection, permeabilized, and stained with rhodamine-phalloidin. Bars, 30 μm. (C) Time-lapse video microscopy was undertaken for HEK293 cells expressing GFP-Nrbl (1-1095) or GFP-Nrbl (1-552) grown on fibronectin-coated coverslips. Images are shown at 5-min intervals with a white arrow highlighting an extending filopodium. Bars, 10 μm.
FIG. 3. PP1 is required for NrbI (1-552)-induced cytoskeletal rearrangement. (A) Lysates (approximately 5% of input used in immunoprecipitations) of HEK293 cells expressing GFP, GFP-NrbI (1-1095), GFP-NrbI (1-552), or GFP-NrbI (1-552, 4A) were immunoblotted with anti-PP1 and anti-GFP. Cells (10^6) were lysed in RIPA buffer and clarified, and GFP-NrbI was immunoprecipitated with anti-GFP. The immunoprecipitates were subjected to SDS–10% (wt/vol) PAGE and analyzed by immunoblotting for PP1. IP, immunoprecipitation; WB, Western blot. (B) Phosphorylase a phosphatase activity associated with the washed GFP-NrbI immunoprecipitates. Phosphatase activity is shown as nanomoles of [32P]phosphate released by equivalent amounts of immunoprecipitates, quantified by immunoblotting with anti-GFP. The data represent averages of three independent experiments performed in duplicate (shown with standard error bars). (C) Sensitivity of the immunoprecipitated GFP-NrbI (1-552)-bound phosphatase activity to okadaic acid, represented as a percentage of that for the control without okadaic acid. The data represent the averages of three independent experiments and are shown with standard error bars. (D) Fluorescence micrographs of Cos7 cells expressing either GFP-NrbI (1-552) or GFP-NrbI (1-552, 4A). Cells were fixed with paraformaldehyde following transfection, permeabilized, and stained with rhodamine-phalloidin. Bar, 20 μm. (E) HEK293 cells expressing GFP, GFP-NrbI (1-552), or GFP-NrbI (1-552, 4A) treated with or without okadaic acid (1 μM) for 15 min prior to fixation. Bars, 25 μm.
not shown). Treatment of GFP-NrbI (1-552)-expressing HEK293 cells with the phosphatase inhibitor okadaic acid (1 μM) (Fig. 3E) or calyculin A (10 nM) rapidly eliminated the filopodia. The brief (15-min) treatment with okadaic acid or calyculin A had no effect on the morphology of cells expressing GFP or GFP-NrbI (1-552, 4A). This indicated that the loss of PP1 binding or inhibition of phosphatase activity had very similar effects on cell morphology and established the importance of PP1 recruitment for NrbI-induced changes in morphology in both Cos7 and HEK293 cells.

Neurabin is a phosphoprotein in cells. While in vitro studies showed PKA phosphorylation of NrbI (17), the phosphorylation of NrbI in vivo in response to PKA agonists had not been investigated. HEK293 cells expressing GFP-NrbI proteins were metabolically labeled with [32P]orthophosphate, and cells were treated with the PKA agonist Sp-cAMPS or the antagonist Rp-cAMPS. The data are representative of three independent experiments. PP1 activity is shown as a percentage of that for the control without GST-NrbI.

FIG. 4. Phosphorylation of NrbI in HEK293 cells. (A) Cells expressing GFP-NrbI (1-1095), GFP-NrbI (1-552), GFP-NrbI (1-552, 4A), GFP-NrbI (553-1095), GFP-NrbI (286-1095), or GFP-NrbI (286-1095, S461A) were metabolically labeled with [32P]orthophosphoric acid for 120 min at 37°C and subsequently treated for 15 min with a PKA agonist, Sp-cAMPS (100 μM), or an antagonist, Rp-cAMPS (100 μM). Cells were lysed in RIPA buffer and clarified, and GFP-NrbI was immunoprecipitated with anti-GFP. Immunoprecipitates were subjected to SDS–7.5% (wt/vol) PAGE, and the gels were dried and autoradiographed. The bottom panel shows relative levels of each GFP-NrbI protein in the immunoprecipitates by immunoblotting with anti-GFP. IP, immunoprecipitation; WB, Western blot. (B) In vitro phosphorylation of His-NrbI (1-615) and His-NrbI (103-615) by PKA and [γ-32P]ATP. The data are representative of three independent experiments. (C) Inhibition of PP1 activity with phosphorylase a as substrate in the presence of increasing concentrations of WT GST-NrbI (374-516) and the mutant S461E. The data are representative of three independent experiments. PP1 activity is shown as a percentage of that for the control without GST-NrbI.
Rapid growth of filopodia is observed in dendrites during the development of the nervous system and following increased synaptic activity. While some filopodia subsequently retract, others proceed to form new spines and synapses. In adult neurons, even mature spines undergo dynamic changes in shape that are mediated by actin rearrangement, and pharmacological agents that promote actin depolymerization result in serious deficits in synaptic efficacy (15). We have analyzed NrbI-bound PP1 as a potential convergence point for signals that regulate both synaptic plasticity and spine morphology. As in vitro studies of NrbI are limited by the inability to express full-length NrbI (4, 17, 20), a key goal of our studies was to establish a cell-based assay that exploited in vitro structure-function analyses of NrbI to yield new insights into the physiological role of NrbI in dictating neuronal morphology, we overexpressed NrbI in 7-DIV-cultured hippocampal neurons, to assess the relevance of NrbI-bound PP1 as a potential convergence point for signals that regulate both synaptic plasticity and spine morphology. As in vitro studies of NrbI are limited by the inability to express full-length NrbI (4, 17, 20), a key goal of our studies was to establish a cell-based assay that exploited in vitro structure-function analyses of NrbI to yield new insights into the physiological role of NrbI in dictating neuronal morphology.
ological role of the NrbI/PP1 complex in the mammalian nervous system.

WT GFP-NrbI (1-1095) bound polymerized F-actin both in vitro and in vivo and modified the morphology of HEK293 and Cos7 cells to reorganize the stress fiber network. GFP-NrbI (286-1095), which lacked actin binding, was cytosolic and had little effect on cell morphology, thereby establishing that actin binding was critical for cytoskeletal reorganization by NrbI. Previous studies showed that NrbI promoted in vitro bundling or cross-linking of actin fibers (20). As ectopically expressed NrbI and NrbII formed homodimers and heterodimers (16), this raised the possibility that the presentation of two adjacent actin-binding domains in a dimeric NrbI may promote actin bundling. Thus, we deleted the C-terminal coiled-coil and SAM domains, shown elsewhere for other proteins to self-associate (14, 27), and by using pull-downs of WT NrbI and NrbII from rat brain extracts established that homodimerization and heterodimerization of neurabin s are mediated via their C termini. The monomeric GFP-NrbI (1-552) induced dramatic changes in cell morphology, disrupting the stress fiber network and promoting highly extended filopodia. Live video microscopy showed the rapid growth of filopodia in the GFP-NrbI (1-552)-expressing cells that extended significantly further than those in cells expressing WT GFP-NrbI (1-1095). The NrbI C terminus (amino acids 553 to 1095), while sufficient to bind NrbI (and NrbII), did not localize with the actin cytoskeleton or modify cell morphology. Cell fractionation and immunoprecipitation studies suggested that GFP-NrbI (1-552) bound both F-actin and PPI more effectively than did WT GFP-NrbI (1-1095), hinting that NrbI dimerization may attenuate its actin- and PPI-binding properties.

The importance of PPI recruitment for NrbI-induced mor-

![Image](http://mcb.asm.org/)
PP1 targeting to cytoskeleton

Phosphorylation was established by mutating the core PP1-binding sequence, KIKF, in NrbI to alanines. This not only abolished PP1 binding in vitro (data not shown) and in vivo, but unlike GFP-NrbI (1-552), GFP-NrbI (1-552, 4A) failed to induce filopodia in HEK293 and Cos7 cells or disrupt stress fibers in Cos7 cells. Pharmacological inhibition of PP1 activity by okadaic acid or calyculin A rapidly collapsed filopodia induced by GFP-NrbI (1-552), showing that, in addition to binding F-actin, NrbI must recruit an active PP1 to modify cell morphology. Immune complex assays confirmed that the NrbI-bound PP1 was an active protein phosphatase. This was unexpected, as in vitro studies (26, 27) (Fig. 4C) showed that PP1-binding NrbI peptides inhibit its activity against phosphorylase a. The molecular basis for the difference in the NrbI/PP1 complexes assembled in vitro and in vivo is unclear, but the difference may reflect the presence of additional proteins or covalent modifications of NrbI in cells. In any case, our data supported an active role for the NrbI/PP1 complex in cytoskeletal reorganization.

PP1 binds both substrates (8, 10) and regulators (22) that undergo reversible phosphorylation. Covalent modification of PP1 regulators modulates PP1 binding (7) and activity (12). Analysis of metabolically labeled HEK293 cells established for the first time that GFP-NrbI (1-1095) was a phosphoprotein and that its phosphorylation in vivo was stimulated by the PKA agonist Sp-cAMPS. Surprisingly, GFP-NrbI (1-552) was essentially unphosphorylated in HEK293 cells. This was not due to elimination of PKA sites, as GFP-NrbI (553-1095) also showed low basal phosphorylation that was not further stimulated by Sp-cAMPS. Deletion of PP1 binding or inhibition of phosphatase activity failed to enhance GFP-NrbI (1-552) phosphorylation, suggesting that PP1 binding did not dictate the phosphorylation state of NrbI. Compared to WT GFP-NrbI (1-1095), soluble GFP-NrbI (286-1095) incorporated much higher levels of [32P]phosphate that was greatly enhanced by PKA activation. This raised the possibility that the actin-binding domain and/or association with the cytoskeleton inhibited NrbI phosphorylation by PKA. This was confirmed by in vitro studies that showed that NrbI (103-615) lacking N-terminal actin-binding sequences incorporated twofold-higher levels of phosphate than did NrbI (1-615). This did not, however, fully explain the absence of phosphate in GFP-NrbI (1-552), leading us to speculate that F-actin association may further impair NrbI phosphorylation in cells. Mutation of serine-461 to alanine in GFP-NrbI (286-1095) abolished PKA-stimulated phosphorylation and identified the serine immediately adjacent to the PP1-binding motif as a major in vivo PKA phosphorylation site. Analysis of recombinant NrbI (374-516, S461E), in which glutamic acid was introduced to mimic serine-461 phosphorylation, confirmed an earlier observation (17) that serine-461 modification diminished PP1 association with NrbI and suggested that assembly-disassembly of the NrbI/PP1 complex may be regulated by hormones that elevate intracellular cyclic AMP.

The PDZ domain adjacent to the core PP1-binding site recruits p70S6K, which has effects opposite of those of PP1 on actin dynamics (23). Affinity isolation of NrbI/PP1 complexes from rat brain or NrbI immunoprecipitation from HEK293 cells overexpressing p70S6K failed to show significant association of p70S6K with NrbI. This demonstrated that NrbI had a significant preference for PP1 over p70S6K and predicted that disruption of PP1 binding would facilitate p70S6K recruitment. Indeed, GFP-NrbI (1-552, 4A) that failed to bind PP1 showed enhanced p70S6K binding. These data can be summarized in the following model for the potential role of NrbI in transducing signals that regulate actin cytoskeleton (Fig. 7). Our data suggest that the majority of cellular NrbI is bound to PP1 and is associated with the actin cytoskeleton. This promotes the disassembly of stress fibers. While both dimeric and monomeric forms of NrbI induce filopodia in cultured cells, the enhanced ability by cells expressing monomeric NrbI to extend surface projections leads us to speculate that the dynamics of NrbI dimerization may also play a role in dictating the cell morphology. Physiological signals that activate Rac1 GTPase (29) promote NrbI shuttling to the cytoskeleton and may target PP1 to the actin cytoskeleton. In contrast, the cytosolic NrbI, by virtue of being an improved substrate for PKA (at serine-461), may display decreased PP1 binding and recruit p70S6K and other targets to the NrbI PDZ domain. This mechanism may also account for the antagonism between PP1 and p70S6K that regulates neuronal morphology. As neither p70S6K overexpression nor inhibition of p70S6K activity by rapamycin had any effect on HEK293 cell morphology, we were unable to establish a role for the NrbI-bound p70S6K in heterologous cells.

NrbI is expressed exclusively in neurons, while NrbII is more...
widely expressed. Both neurabin1s and neurabin2 are retained in a complex with the phosphatase type 1 catalytic subunit. Genes Dev. 10:555–569.


