

# $\delta$ -Catenin-induced Dendritic Morphogenesis

## AN ESSENTIAL ROLE OF p190RhoGEF INTERACTION THROUGH AKT1-MEDIATED PHOSPHORYLATION<sup>\*,§</sup>

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$\delta$ -Catenin was first identified through its interaction with Presenilin-1 and has been implicated in the regulation of dendrogenesis and cognitive function. However, the molecular mechanisms by which  $\delta$ -catenin promotes dendritic morphogenesis were unclear. In this study, we demonstrated  $\delta$ -catenin interaction with p190RhoGEF, and the importance of Akt1-mediated phosphorylation at Thr-454 residue of  $\delta$ -catenin in this interaction. We have also found that  $\delta$ -catenin overexpression decreased the binding between p190RhoGEF and RhoA, and significantly lowered the levels of GTP-RhoA but not those of GTP-Rac1 and -Cdc42.  $\delta$ -Catenin T454A, a defective form in p190RhoGEF binding, did not decrease the binding between p190RhoGEF and RhoA.  $\delta$ -Catenin T454A also did not lower GTP-RhoA levels and failed to induce dendrite-like process formation in NIH 3T3 fibroblasts. Furthermore,  $\delta$ -catenin T454A significantly reduced the length and number of mature mushroom shaped spines in primary hippocampal neurons. These results highlight signaling events in the regulation of  $\delta$ -catenin-induced dendrogenesis and spine morphogenesis.

$\delta$ -Catenin was first identified by yeast two-hybrid screening as a molecule that interacts with Presenilin-1 (PS-1),<sup>3</sup> which is the most prominently mutated gene in familial Alzheimer disease (FAD) patients (1, 2). The interaction of  $\delta$ -catenin with PS-1, along with its abundant expression in neurons, suggests

that  $\delta$ -catenin has specialized neuronal functions (3, 4). Indeed,  $\delta$ -catenin-deficient mice showed severe learning deficits and abnormal synaptic plasticity, suggesting a special role of  $\delta$ -catenin at the synapse (5). Furthermore, the hemizygous loss of the chromosomal 5p15.2 region, which contains the human  $\delta$ -catenin gene, results in the severe mental retardation associated with Cri du Chat syndrome. This chromosomal abnormality may account for 1% of all mentally retarded individuals (6).

Structural analysis indicated that  $\delta$ -catenin is a member of the p120-Catenin (hereafter, p120<sup>ctn</sup>) subfamily of armadillo proteins and has a DS<sub>WV</sub> sequence at the C terminus that binds to the PDZ (PSD-95/Disc-large/ZO-1) domain-containing proteins (7).  $\delta$ -Catenin also contains SH3 binding domains at the N terminus (4, 8), a GKKKKKKK sequence (putative NLS) that can potentially promote lipid intermixing (9), and a proline-rich domain that is likely to be involved in the interaction with the actin-binding protein, Profilin (4). The presence of 10 Arm repeats in  $\delta$ -catenin suggests its potential participation in various protein-protein interactions. In addition to PS-1, the  $\delta$ -catenin-associated proteins identified thus far include E-cadherin (4), S-SCAM (7), p0071 (10), Densin-180 (11), PSD-95, Abl (8), Cortactin (12), sphingosine kinase (13), and Kaiso (14), suggesting its many possible roles in cells. Our previous reports demonstrated that the overexpression of  $\delta$ -catenin induces the branching of dendrite-like processes in both NIH 3T3 fibroblasts and primary hippocampal neurons (15). We have also reported that an E18 hippocampal neuron overexpressing  $\delta$ -catenin demonstrates enhanced arborization of dendrites, swelling, and maturation of dendritic spines (15, 16).  $\delta$ -Catenin-induced dendrogenesis is quite similar in shape to the morphological changes induced by treatment with C3 exotoxin, which is a specific Rho inhibitor derived from *Clostridium botulinum* (12, 17). The inhibition of Rho signaling also enhances not only the formation of the p120<sup>ctn</sup>-induced branch (18) but also  $\delta$ -catenin-induced dendrogenesis (12). However, the underlying mechanism for how  $\delta$ -catenin overexpression affects Rho signaling is unclear.

The Rho family of small GTPases, consisting of Cdc42, Rac, and Rho, has been implicated in the control of numerous cellular processes including cytoskeletal reorganization, transcriptional activation, cell cycle progression, cell fate determination, and synaptic plasticity (17, 19, 20). Of the Rho GTPase family members, Cdc42, Rac1, and RhoA have been studied most

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<sup>3</sup> The abbreviations used are: PS-1, Presenilin-1; FL, full-length; GEF, guanine nucleotide exchange factor; GFP, green fluorescent protein; MEF, mouse embryonic fibroblast; wt, wild type; ANOVA, analysis of variance; HA, hemagglutinin; GST, glutathione S-transferase.

extensively. The activation of the Rho family proteins requires GDP-GTP exchange that is catalyzed by various guanine nucleotide exchange factors (GEFs), and the cessation of their actions is regulated by GTPase-activating proteins (GAPs) and guanine nucleotide dissociation inhibitors (GDIs) (21–23). Among the Rho-specific GEFs, p190RhoGEF was first identified as a brain-enriched GDP/GTP exchange factor that activates RhoA and contains the predicted  $\alpha$ -helical coiled-coil domain in the C-terminal (24, 25). The overexpression of p190RhoGEF mimics activated RhoA in stimulating cytoskeletal contraction and preventing neurite outgrowth (25). The C-terminal of p190RhoGEF has been shown to interact with microtubules (25), JIP-1 (26), a destabilizing element in the 3'-UTR of neurofilament light (NF-L) mRNA (27), FAK (focal adhesion kinase) (28), and 14-3-3 (29), which is important for its regulatory function in cells. Furthermore, activation of Rho GTPases by p190RhoGEF leads to a decrease in dendritic branching, whereas GDP-Rho (inactive) induces dendrogenesis (24). Among the binding partners of p190RhoGEF, 14-3-3 was recently identified as a new binding partner of  $\delta$ -catenin as well (30). Most 14-3-3 binds to the phosphoserine/threonine residue on a partner protein when it is phosphorylated by various kinases including protein kinase A, protein kinase C, and Akt (31–33). Akt/PKB, a serine/threonine protein kinase, regulates multiple biological processes including synaptic strength, cell survival, proliferation, growth, and glycogen metabolism (34–36). In mammals, Akt/PKB is composed of three Akt family members such as Akt1/PKB $\alpha$ , Akt2/PKB $\beta$ , and Akt3/PKB $\gamma$ , and is activated by various growth and survival factors through a pathway requiring PI3K-dependent generation of phosphatidylinositol (3, 4, 5)trisphosphate (37, 38).

In this study, we found that  $\delta$ -catenin interacts with p190RhoGEF and significantly lowered the levels of GTP-RhoA, for which Akt1 plays an important role through the Thr-454 phosphorylation on  $\delta$ -catenin. By introducing a substitution of Thr-454 residue on  $\delta$ -catenin to Ala, we demonstrated that  $\delta$ -catenin T454A, a defective form in binding p190RhoGEF, significantly reduced the length and number of mature mushroom shaped spines in primary hippocampal neurons. Our results suggest the interaction between  $\delta$ -catenin and p190RhoGEF can act as a key modulator in regulating  $\delta$ -catenin-induced dendrogenesis and spine formation.

## EXPERIMENTAL PROCEDURES

**Plasmids and Antibodies**—The construction of  $\delta$ -catenin full-length (FL-) and  $\Delta$ C207-in pEGFP-C1 has been previously described (15). The  $\delta$ -catenin T454A mutant, in which the Thr-454 residue was substituted to Ala, was generated by site-directed mutagenesis. The expression plasmid for p190RhoGEF (pcDNA3-HA) was kindly provided by W. Moolenaar (The Netherlands Cancer Institute, Amsterdam, The Netherlands); Myc-RhoA by K. Y. Lee (Chonnam National University, Gwanju, Korea).

The antibodies were obtained as follows: anti- $\delta$ -catenin (07-259) and anti-Rac1 (05-389) (Upstate Biotechnology); anti-GFP (632376), anti- $\delta$ -catenin (C98320) (BD Biosciences); anti- $\beta$ -tubulin and anti-phosphothreonine (P6623, Sigma); and anti-RhoA (sc-418n), anti-Cdc42 (sc-87), and anti-phosphoserine

(sc-57555, Santa Cruz Biotechnology). The antibody for p190RhoGEF was kindly provided by D. Schlaepfer and B. Margolis. HA or Myc epitopes were detected using media from 12CA5 or 9E10 hybridoma, respectively.

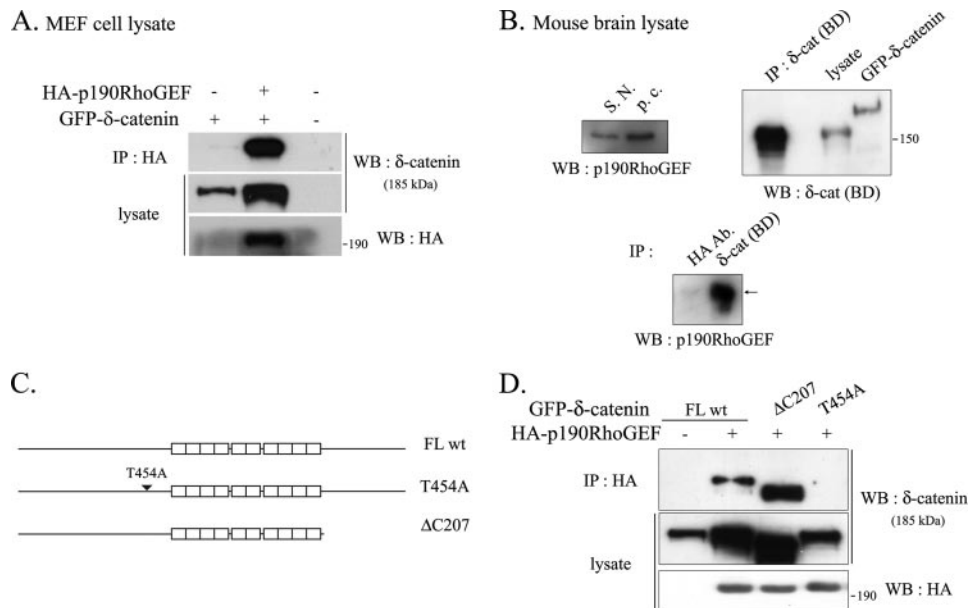
**Cell Culture and Transfection**—The NIH 3T3, mouse embryonic fibroblast (MEF), and Bosc23 cells were grown in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum and 1% penicillin/streptomycin at 37 °C with 5% CO<sub>2</sub>. The cells were transfected using calcium phosphate or Lipofectamine Plus reagent (Invitrogen), according to the manufacturer's instructions.

**Immunoblotting and Akt Kinase Assay**—Immunoprecipitation/immunoblotting was performed as previously described (39). For the kinase assays, the full-length and mutant GFP- $\delta$ -catenin constructs were transiently overexpressed in MEF cells or Bosc23 cells. To collect P1 mouse brain lysates, 1 day postnatal mouse were sacrificed.  $\delta$ -Catenin proteins were purified by immunoprecipitation using the specific antibody and protein G-Sepharose (BD Biosciences). The immune complexes were washed three times with lysis buffer and twice with kinase buffer, and then incubated for 30 min at 30 °C in a 20- $\mu$ l reaction mixture supplemented with recombinant Akt protein and either 10  $\mu$ M cold ATP or 10  $\mu$ Ci of [ $\gamma$ -<sup>32</sup>P]ATP. Akt, cold ATP, and kinase buffer were purchased from Cell Signaling Technology and [ $\gamma$ -<sup>32</sup>P]ATP was purchased from BMS. Radioactive phosphorylation status of  $\delta$ -catenin was detected by autoradiograph and nonradioactive phosphorylation status was determined by immunoblotting using the phospho-Ser or phospho-Thr antibodies. Equal loading of proteins was confirmed using EZ staining kits (EZBioPaQ, Co., Ltd.).

**Affinity Precipitation of Cellular GTPases**—The cellular RhoA activity was determined using GST-RBD as previously described by Ren *et al.* (40). The cells were lysed in lysis buffer (50 mM Tris, pH 7.4, 1% Triton X-100, 0.5% sodium deoxycholate, 0.1% SDS, 500 mM NaCl, 10 mM MgCl<sub>2</sub>, and protease inhibitor mixture). The lysates were incubated with the GST-RBD beads (Cytoskeleton Inc.) at 4 °C for 1 h. The beads were then washed four times with washing buffer (50 mM Tris, pH 7.4, 1% Triton X-100, 150 mM NaCl, 10 mM MgCl<sub>2</sub>, and protease inhibitor mixture). The bound RhoA proteins were detected by immunoblotting using a monoclonal antibody against RhoA.

The cellular Rac1 and Cdc42 activities were determined using GST-PBD as described in other studies (41). The cDNA of the p21-binding domain (PBD) from PAK1 (amino acids 67–150) was cloned into the bacterial expression vector pGEX-4T3 and expressed as a GST fusion protein. The purified fusion proteins were isolated from glutathione-Sepharose 4B beads. The lysates were incubated with the GST-PBD beads, and the beads were washed twice with 25 mM Tris, pH 7.4, 1 mM dithiothreitol, 30 mM MgCl<sub>2</sub>, 40 mM NaCl, 1% Nonidet P-40, and twice with the same buffer without Nonidet P-40. The bound Rac1 and Cdc42 proteins were detected by immunoblotting using their specific antibodies.

The relative activity of each GTPase was determined by quantifying each band of GTP-bound GTPase and the total amount of GTPase using the TINA 2.09 software program (Raytest), and the values of the GTP-bound bands were normalized to the value of the total amount. All results were deter-



**FIGURE 1.  $\delta$ -Catenin interacts with p190RhoGEF, for which Thr-454 residue of  $\delta$ -catenin is indispensable.** A, MEF cells were transfected with GFP-tagged  $\delta$ -catenin with or without HA-p190RhoGEF. The binding of  $\delta$ -catenin and p190RhoGEF was examined by immunoprecipitation with the anti-HA antibody, and Western blotting was performed with the anti- $\delta$ -catenin antibody (top panel). The expression of  $\delta$ -catenin or p190RhoGEF in the cell lysates were detected using the anti- $\delta$ -catenin antibody (middle panel), or the anti-HA antibody (bottom panel), respectively. B, adult mouse brain lysates were used to examine the endogenous interaction between  $\delta$ -catenin and p190RhoGEF (bottom panel). HA antibody was used as a negative control for immunoprecipitation assay. Endogenous expression of p190RhoGEF in the adult mouse brain region of substantia nigra (indicated as S.N.) and pars compacta (indicated as p.c.) were shown (upper left panel). Endogenous expression of  $\delta$ -catenin in the whole adult mouse brain lysate was shown, and the efficiency of immunoprecipitation using the anti- $\delta$ -catenin antibody was determined with the same adult mouse brain lysates (upper right panel). C, schematic illustration of  $\delta$ -catenin constructs used throughout experiments. D, MEF cells were transfected with either GFP- $\delta$ -catenin FL wt or a mutant of  $\delta$ -catenin ( $\Delta$ C207 and T454A) together with or without HA-p190RhoGEF as indicated in the figure. The binding of  $\delta$ -catenin and p190RhoGEF was examined by immunoprecipitation with the anti-HA antibody, and blotted with the anti- $\delta$ -catenin antibody (upper panel). The expression of  $\delta$ -catenin or p190RhoGEF in the cell lysates was detected with the anti- $\delta$ -catenin antibody (middle panel) or the anti-HA antibody (bottom panel), respectively.

mined using three different film exposures from at least three independent experiments.

**Hippocampal Neuron Culture and Transfection**—Cultured hippocampal neurons were prepared from embryonic day 18 (E-18) fetal Sprague-Dawley rats and plated on poly-D-lysine coated five 18-mm glass coverslips at a density of 200,000 cells/60-mm dish. The cultures were grown in Neurobasal medium (Invitrogen) supplemented with 2% B-27, and 0.5 mM L-glutamine. The neurons were transfected at 16 DIV using the calcium-phosphate-mediated method (42). Briefly, 8–10  $\mu$ g of cDNA and 7.5  $\mu$ l of 2 M  $\text{CaCl}_2$  were mixed in distilled water to a total volume of 75  $\mu$ l, and same volume of 2 $\times$  BBS was added. The cell culture medium was completely replaced by transfection medium (MEM, 1 mM pyruvate, 0.6% glucose, 10 mM glutamine, and 10 mM HEPES, pH 7.65), and the cDNA mixture was added to the cells, which were then incubated in a 5%  $\text{CO}_2$  incubator for 90 min. They were washed twice with transfection medium (pH 7.35) and then returned to the original culture medium. Transfection efficiency was about 1–5%.

**Image Acquisition and Data Analysis**—The cells were fixed in 4% paraformaldehyde/4% sucrose/PBS for 15 min, washed 2  $\times$  5 min, and permeabilized for 5 min in 0.25% Triton X-100/phosphate-buffered saline. The images were obtained using an Olympus IX71 microscope (Olympus) with 40 $\times$  N.A. 1.0 or

60 $\times$  N.A. 1.4 oil lens using a CoolSNAP-Hq CCD camera (Roper Scientific) driven by MetaMorph imaging software (Universal Imaging Co.). Light from a mercury lamp was shuttered using a VMM1 Uniblitz shutter (Vincent Associates). The number of dendritic branches was analyzed with a sphere, 100  $\mu$ m in radius, centered at the soma. The number of intersections between the dendritic branches and a sphere was averaged. Statistical analysis was performed using ANOVA and Tukey's HSD post hoc test. The analysis and quantification of the data were performed using MetaMorph software and SigmaPlot 8.0 (Systat Software). The expression of each construct was confirmed by the retrospective immunostaining using specific antibodies, and only immunopositive neurons were included in the analysis. Data analysis was performed in a blinded manner. The data are presented as the mean  $\pm$  S.E.

## RESULTS

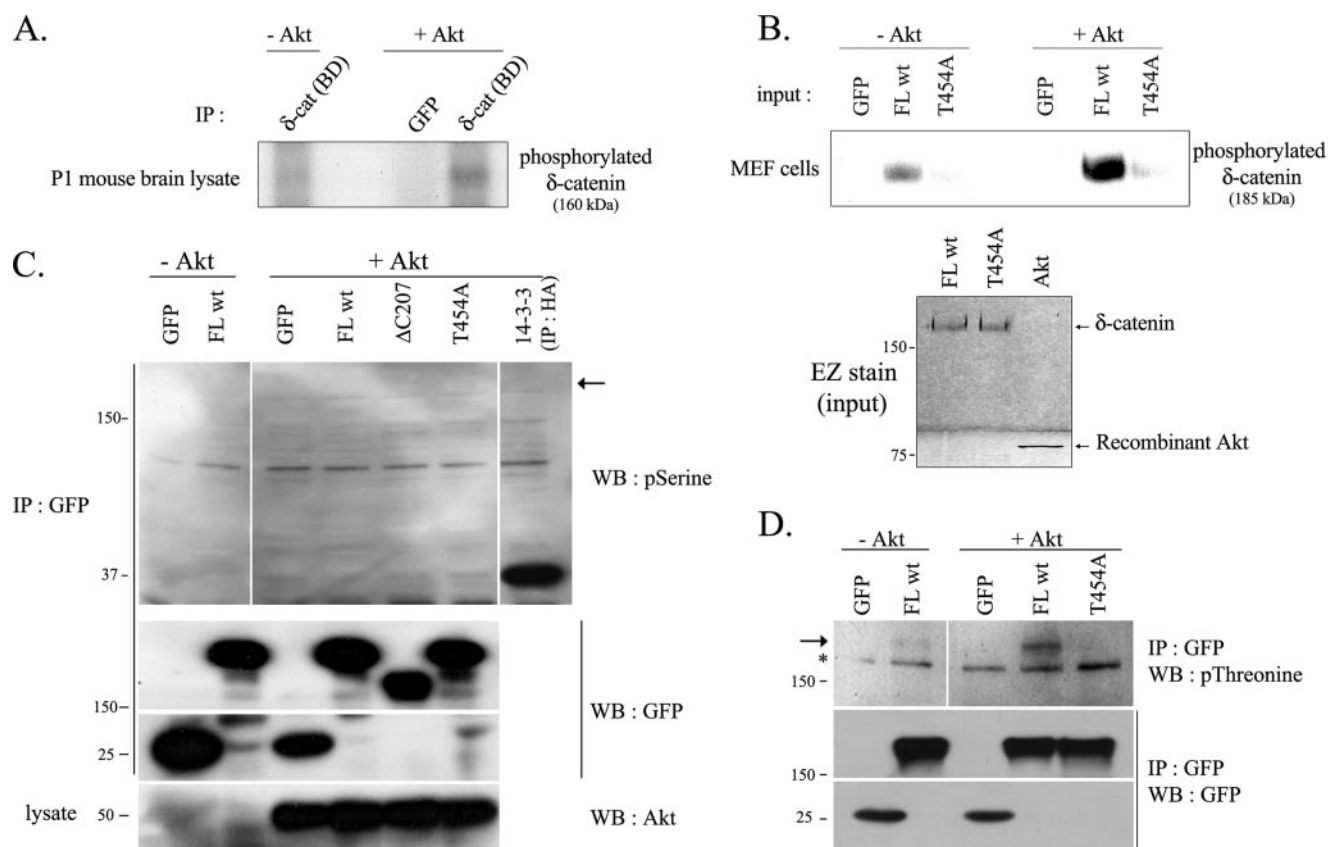
**$\delta$ -Catenin Interacts with p190RhoGEF, for Which the Thr-454 Residue of  $\delta$ -Catenin Is Indispensable**—The published results have shown that 14-3-3 can interact with either

p190RhoGEF (29) or  $\delta$ -catenin (30), but association between p190RhoGEF and  $\delta$ -catenin has not been demonstrated until now. Therefore, the interaction between  $\delta$ -catenin and p190RhoGEF was examined by co-transfecting HA-tagged p190RhoGEF into wild-type MEFs with GFP-tagged  $\delta$ -catenin, followed by immunoprecipitation analysis. The results indicated that  $\delta$ -catenin specifically binds to the p190RhoGEF (Fig. 1A). The interaction between  $\delta$ -catenin and p190RhoGEF was also confirmed using endogenous proteins. As shown in Fig. 1B, mouse brain expressed both endogenous  $\delta$ -catenin and p190RhoGEF, and endogenous p190RhoGEF was specifically co-immunoprecipitated with  $\delta$ -catenin. To map the region responsible for the interaction between  $\delta$ -catenin and p190RhoGEF, we used a site-directed  $\delta$ -catenin mutant and a deletion mutant as shown in a schematic diagram in Fig. 1C. The immunoprecipitation assay showed that p190RhoGEF could interact with  $\delta$ -catenin FL wt and  $\Delta$ C207 but not with the T454A mutant (Fig. 1D).

Our protein motif scan analysis revealed that mouse  $\delta$ -catenin contains 3 putative 14-3-3 binding sites (Ser-282, Thr-454, Ser-1094). We thus investigated the possibility of 14-3-3 requirement for the interaction between  $\delta$ -catenin and p190RhoGEF. The deletion constructs were specifically designed to remove the putative 14-3-3 binding residue on



## Effects of $\delta$ -Catenin on Dendrogenesis and Spine Formation

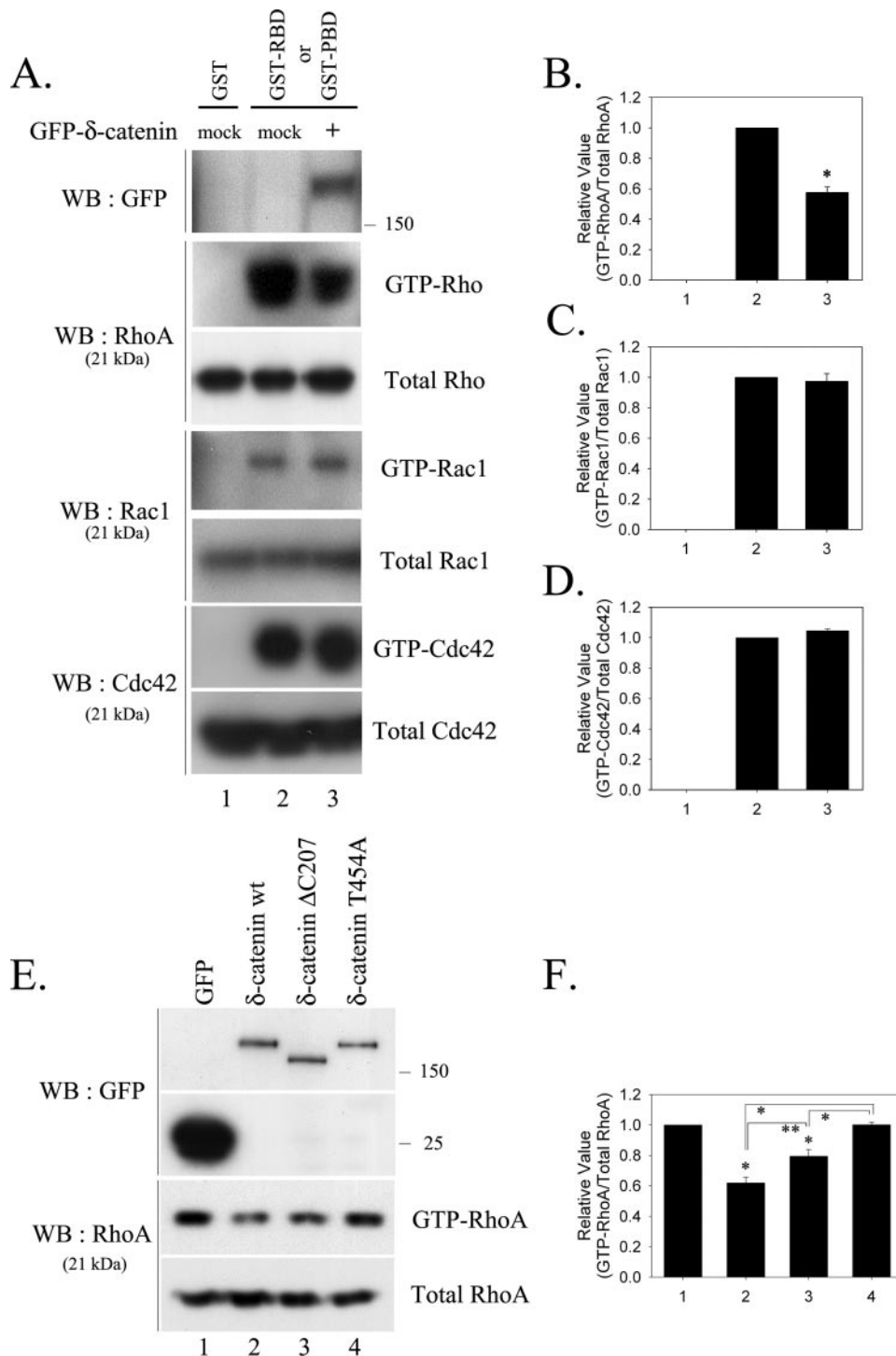


**FIGURE 2.  $\delta$ -Catenin undergoes Akt1-mediated phosphorylation at Thr-454 residue.** An Akt1 kinase assay was performed as described under "Experimental Procedures." GFP, mock-transfected proteins were used as a negative control in the assay, and GFP-immunocomplex was obtained by immunoprecipitating with GFP antibody. **A**, purified immunocomplex of 1 day post-natal mouse brain lysates with  $\delta$ -catenin (BD) antibody was used as a substrate for an Akt kinase assay. [ $\gamma$ - $^{32}$ P]ATP was supplemented with a reaction buffer, and phosphorylation status was detected by autoradiograph. **B**, GFP-tagged  $\delta$ -catenin wt (FL wt) and T454A mutant were transfected in MEF cells, and purified immunocomplex with  $\delta$ -catenin (BD) antibody was used as a substrate for an Akt kinase assay. [ $\gamma$ - $^{32}$ P]ATP was supplemented with a reaction buffer, and phosphorylation status was detected by autoradiograph (upper panel). The level of  $\delta$ -catenin FL wt and T454A mutant existing in immunocomplex and of recombinant Akt1 were confirmed by EZ staining kit (bottom panel). **C** and **D**, GFP-tagged full-length (FL wt) and mutant  $\delta$ -catenin ( $\Delta$ C207 and T454A) were overexpressed in Bosc23 cells, and purified immunocomplex with GFP antibody was used as a substrate for Akt kinase assay. Radio-inactive cold ATP was supplemented with a reaction buffer, and phosphorylation status on serine and threonine residue in the wild type or mutant type  $\delta$ -catenin was detected using anti-phosphoserine antibody (**C**) or with anti-phosphothreonine antibody (**D**) (upper panel). HA-14-3-3 was immunoprecipitated with HA antibody and was used as a positive control (**C**). An arrow indicates the predicted site (**C**) or an actual band (**D**) of phosphorylated  $\delta$ -catenin, and an asterisk indicates a nonspecific band. The immunocomplexes were subsequently reprobated with anti-GFP antibody (middle panels), and the added recombinant Akt1 was also confirmed using anti-Akt antibody (**C**, bottom panel).

$\delta$ -catenin as follows:  $\Delta$ N85-325-(deletion of Ser-282),  $\Delta$ C207-(deletion of Ser-1094), site-directed mutant,  $\delta$ -catenin T454A. As shown in Supplemental Fig. S1A, all of these  $\delta$ -catenin deletion mutants were specifically immunoprecipitated by 14-3-3 $\epsilon$ , but  $\delta$ -catenin  $\Delta$ N85-325 bound to 14-3-3 $\epsilon$  to a lesser extent (Supplemental Fig. S1A). Interestingly, FL wt,  $\Delta$ N85-325, and  $\delta$ -catenin T454A had a noticeable interaction with 14-3-3 $\zeta$ , whereas  $\delta$ -catenin  $\Delta$ C207 barely interacted (Supplemental Fig. S1B), suggesting a 14-3-3 isoform-specific interaction with  $\delta$ -catenin and 14-3-3. Exceptionally,  $\delta$ -catenin T454A, a defective form in p190RhoGEF binding, retained the capability to bind both 14-3-3 isoforms, suggesting that 14-3-3 does not mediate the interaction between  $\delta$ -catenin and p190RhoGEF.

**$\delta$ -Catenin Undergoes Akt1-mediated Phosphorylation at Thr-454 Residue**—As our protein motif scan analysis showed that the Thr-454 residue in mouse  $\delta$ -catenin could also be a putative target phosphorylation site for Akt, we investigated the possibility of Akt1-mediated  $\delta$ -catenin phosphorylation. An *in vitro* Akt1 kinase assay was performed using endogenous  $\delta$ -catenin in 1 day post-natal mouse brain lysates and GFP-

tagged exogenous  $\delta$ -catenin (Fig. 2, **A** and **B**, respectively). As shown in Fig. 2, **A** and **B**, both endogenous and exogenous wild type  $\delta$ -catenin were specifically phosphorylated by Akt1.  $\delta$ -Catenin T454A, however, was not phosphorylated by Akt1, indicating that the Thr-454 residue in mouse  $\delta$ -catenin is a target site for Akt1-phosphorylation. To confirm these results, immunoblotting with specific anti-phospho-Ser or -Thr antibodies was performed (Fig. 2, **C** and **D**, respectively). 14-3-3 $\zeta$ , a well-known target protein that can be phosphorylated by Akt on its Ser-58 residue, was used as a positive control to certify the activity of anti-phospho-Ser antibody, and no phosphorylated  $\delta$ -catenin band by Akt1 was detected (Fig. 2C). This demonstrates that Akt1 is not responsible for phosphorylation of the Ser-282 or Ser-1094 residues in  $\delta$ -catenin. In contrast, phosphorylated  $\delta$ -catenin FL wt, but not T454A mutant band, was detected by antibody against phosphothreonine residues (Fig. 2D). This result again proved the principal role of Thr-454 residue on  $\delta$ -catenin in Akt1-mediated phosphorylation. Akt1 is inferred as an important regulator for p190RhoGEF interaction with  $\delta$ -catenin from a series of experiments. The increased



**FIGURE 3. Overexpression of  $\delta$ -catenin results in a decrease in the level of GTP-RhoA, while the Rac1 and Cdc42 activity are unaffected.** A, levels of active GTP-bound RhoA, Rac1, and Cdc42 were measured in MEF cells transfected with either the mock or GFP- $\delta$ -catenin. The RhoA activity was measured using a GST-RBD, and the Rac1, Cdc42 activities were measured using a GST-PBD. The total amounts of RhoA, Rac1, and Cdc42 are also shown to compare the endogenous level of each GTPase in the transfected cells. The relative activity of RhoA (B), Rac1 (C), and Cdc42 (D) were determined as described under "Experimental Procedures." The data are represented as the mean  $\pm$  S.E. (\*,  $p < 0.01$ ). E, level of active GTP-bound RhoA was measured in the MEF cells transfected with GFP or with wt/mutant GFP- $\delta$ -catenin constructs using GST-RBD. F, relative activity of RhoA was determined. The data are represented as the mean  $\pm$  S.E. (\*,  $p < 0.01$ ; \*\*,  $p < 0.05$ ). Results are representative of at least three independent experiments.

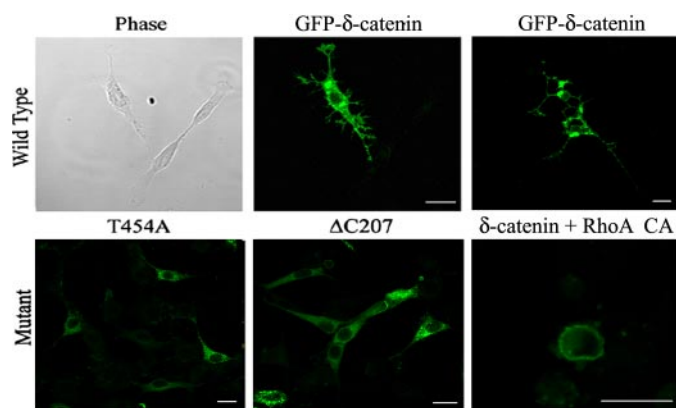
amounts of either wild type or dominant negative p190RhoGEF increased the levels of  $\delta$ -catenin (Supplemental Fig. S2). But only wild-type Akt1 showed dose-dependent increases in the

level of  $\delta$ -catenin, whereas kinase dead form of Akt1 slightly decreased the level of  $\delta$ -catenin (Supplemental Fig. S3). These results suggest that association of p190RhoGEF with  $\delta$ -catenin followed by Akt1-mediated phosphorylation enhances the stability of  $\delta$ -catenin and GEF activity of p190RhoGEF is not a critical factor for the association.

**Overexpression of  $\delta$ -Catenin Significantly Reduces the Level of GTP-RhoA, While the Level of GTP-Rac1 and -Cdc42 Are Not Affected**—Even though there are many reports showing that small GTPase family proteins including Rac1, Cdc42, and RhoA affect dendrogenesis (17, 19), there is no evidence showing how the activities of these proteins are affected by  $\delta$ -catenin overexpression. In this report, we propose a model showing that overexpressed  $\delta$ -catenin can lower GTP-RhoA levels by sequestering p190RhoGEF from its substrate RhoA, thereby attenuating RhoA activation. GST-PBD (p21-binding domain) and GST-RBD (Rho-binding domain) were used, and GST pull-down assays were performed to determine the levels of the GTP-bound Rac1 and Cdc42, and RhoA, respectively, using their specific antibodies.  $\delta$ -Catenin overexpression significantly decreased the level of GTP-RhoA by  $\sim 40\%$  compared with those in the control cells (Fig. 3, A and B). As a comparison, overexpression of  $\delta$ -catenin  $\Delta$ C207 significantly decreased the level of GTP-RhoA by  $\sim 20\%$  compared with those in the GFP-overexpressed control cells.  $\delta$ -Catenin T454A, however, did not affect the level of GTP-RhoA (Fig. 3, E and F). In contrast, the levels of GTP-Rac1 and -Cdc42 were not affected by  $\delta$ -catenin overexpression (Fig. 3, A and C and D).

**Effects of the Wild Type and Mutant  $\delta$ -Catenin on the Dendrite-like Process Formation in NIH 3T3 Fibroblasts and Mature Mushroom-**

**shaped Spine Formation in the Primary Hippocampal Neurons**—Previously, we reported that  $\delta$ -catenin expression induces the branching of dendrite-like processes, and that  $\delta$ -catenin  $\Delta$ C207



**FIGURE 4. Effects of the wild type and mutant  $\delta$ -catenins on the dendrite-like process formation in NIH 3T3 fibroblasts.** The NIH 3T3 fibroblast cells were transfected with wild type (GFP- $\delta$ -catenin) and mutant  $\delta$ -catenin (T454A and  $\Delta$ C207) or co-transfected with a constitutive active mutant of RhoA (RhoA CA) as indicated in each image. After 24 h post-transfection, the cells were fixed, and a fluorescent image was taken sometimes together with its corresponding phase contrast image (left and middle, upper panel). GFP- $\delta$ -catenin images on top panel indicate typical  $\delta$ -catenin-induced dendrite-like processes at different stages (upper middle-early stage; upper right-late stage). Scale bars, 20  $\mu$ m.

does not induce either dendrite-like branching or significant extension of processes in NIH 3T3 fibroblasts and enhances dendritic morphogenesis in primary hippocampal neurons (15). In this study, the ability of  $\delta$ -catenin T454A to induce dendrite-like processes formation in NIH 3T3 cells was examined in comparison with  $\delta$ -catenin full-length and  $\Delta$ C207 mutant. As shown in Fig. 4,  $\delta$ -catenin T454A did not induce any noticeable morphological changes, suggesting that the association of  $\delta$ -catenin with p190RhoGEF is essential for the  $\delta$ -catenin induced dendrite-like process formation in fibroblasts. 100% of wild-type  $\delta$ -catenin overexpressing cells showed dendrite-like process formation in quantitative analysis but neither  $\delta$ -catenin  $\Delta$ C207 nor T454A-overexpressing cells showed any process formation. The co-expression of constitutive active form of RhoA with wild-type  $\delta$ -catenin totally abolished the dendrite-like process formation, and resulted in a rounded morphology ( $\delta$ -catenin + RhoA CA in Fig. 4). This suggests that the RhoA signaling might be a downstream effector of  $\delta$ -catenin.

We have also previously described that E18 hippocampal neurons overexpressing the full-length  $\delta$ -catenin demonstrated enhanced arborization of dendrites, swelling, and mature dendritic spines (15). To examine the effects of  $\delta$ -catenin T454A on dendrogenesis and spine formation,  $\delta$ -catenin and its variants were transfected in cultured hippocampal neurons (Fig. 5). In the  $\delta$ -catenin transfected neurons, the number of mature spines, either a cotyloid appearance or flat-apex mushroom appearance, was significantly higher compared with the GFP control ( $0.38 \pm 0.03/\mu\text{m}$ ,  $n = 10$ , for GFP control;  $0.5 \pm 0.01/\mu\text{m}$ ,  $n = 10$ , for  $\delta$ -catenin,  $n = 18$ ,  $p < 0.01$ ). The expression of  $\delta$ -catenin  $\Delta$ C207 also increased the number of mature spines ( $0.45 \pm 0.02/\mu\text{m}$ ,  $n = 12$ ) but to a significantly lesser extent than the FL wt  $\delta$ -catenin ( $p < 0.05$ ).  $\delta$ -Catenin T454A, which is not able to bind p190RhoGEF, decreased the number of mature spines most significantly compared with the full-length  $\delta$ -catenin or  $\delta$ -catenin  $\Delta$ C207 ( $0.4 \pm 0.04/\mu\text{m}$ ,  $n = 12$ , for  $\delta$ -catenin

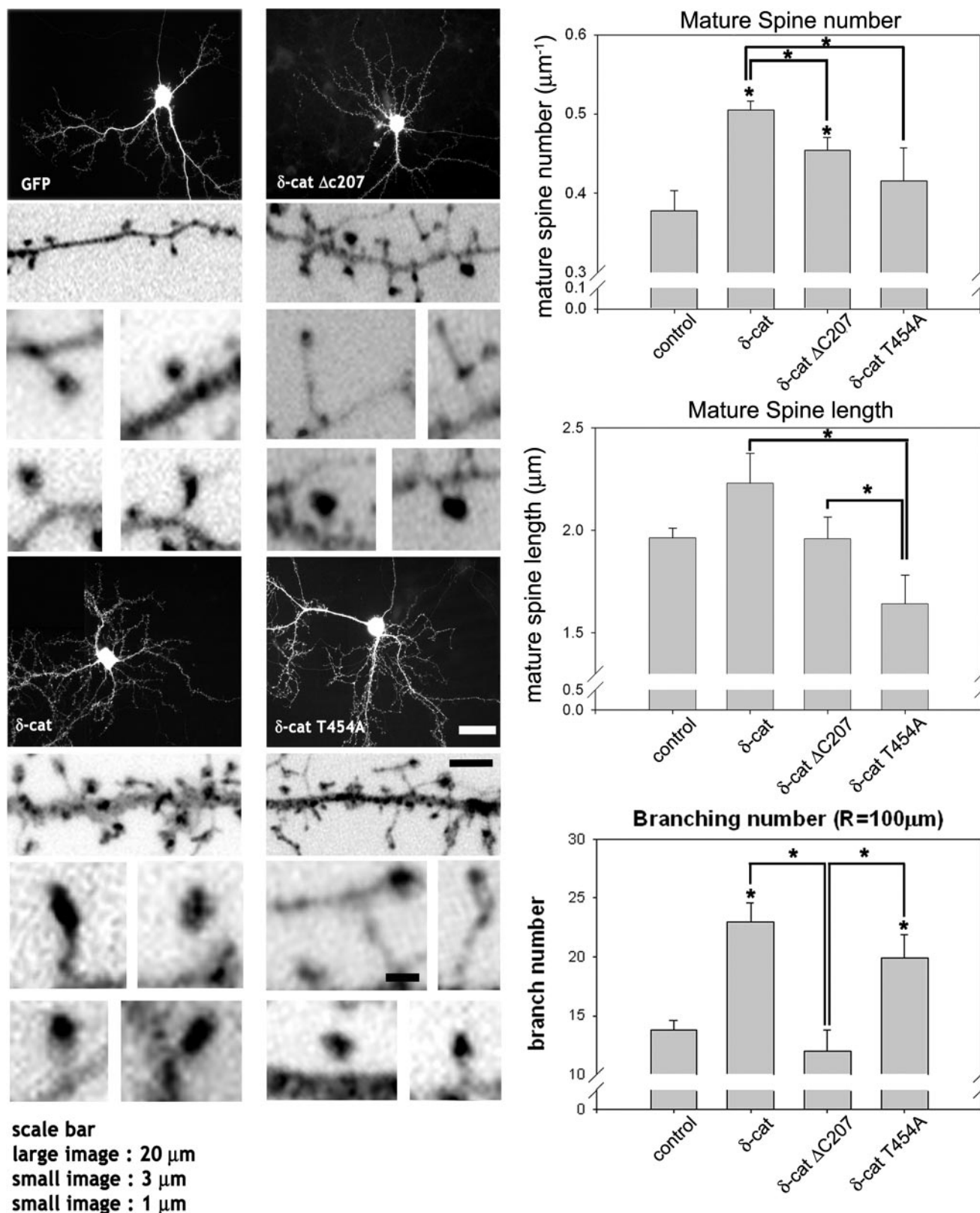
T454A,  $p < 0.01$ ). Surprisingly, the morphology of the dendritic spines in hippocampal neurons overexpressing  $\delta$ -catenin T454A dramatically changed. Typical mushroom shaped spines were rare and most spines showed rather long and thin filopodial morphologies. When only the length of mushroom-shaped spines was compared,  $\delta$ -catenin T454A expression significantly reduced the length of the spines when compared with the full-length  $\delta$ -catenin or its  $\Delta$ C207 mutant ( $1.54 \pm 0.11 \mu\text{m}$ ,  $n = 10$ , for  $\delta$ -catenin T454A;  $2.22 \pm 0.14 \mu\text{m}$ ,  $n = 10$ , for full-length  $\delta$ -catenin;  $1.95 \pm 0.10 \mu\text{m}$ ,  $n = 10$ , for  $\delta$ -catenin  $\Delta$ C207,  $p < 0.01$ ). Consistent with a previous report (15), the expression of  $\delta$ -catenin FL also increased the branching of dendrites ( $13.8 \pm 0.8$ ,  $n = 12$ , for GFP control;  $23.2 \pm 1.5$ ,  $n = 12$ , for  $\delta$ -catenin FL,  $p < 0.01$ ). Unexpectedly,  $\delta$ -catenin T454A also increased the number of dendrite branches while its  $\Delta$ C207 mutant did not ( $19.9 \pm 2$ ,  $n = 10$ , for  $\delta$ -catenin T454A,  $p < 0.01$  compared with GFP control;  $10.9 \pm 0.7$ ,  $n = 10$ , for  $\delta$ -catenin  $\Delta$ C207).

**$\delta$ -Catenin Sequesters p190RhoGEF from RhoA**—As shown in Fig. 6, we proposed a model illustrating Akt1-mediated Thr-454 phosphorylation on  $\delta$ -catenin leads to its interaction with p190RhoGEF, which sequesters p190RhoGEF from activating Rho signaling. Thereby, the sequestration of p190RhoGEF by  $\delta$ -catenin contributes to the attenuated Rho signaling, resulting in enhanced dendrogenesis and spine maturation. To prove our proposed model, we performed the immunoprecipitation assays with  $\delta$ -catenin, p190RhoGEF, and RhoA. If the interaction between  $\delta$ -catenin and p190RhoGEF indeed sequesters p190RhoGEF from binding to RhoA, the  $\delta$ -catenin overexpression should decrease the interaction between p190RhoGEF and RhoA. As  $\delta$ -catenin is known to interact with a cytoplasmic loop of Presenilin-1, HA-tagged Presenilin-1 was used as a positive control for  $\delta$ -catenin interaction, and as Notch is a well-known substrate for Presenilin-1/ $\gamma$ -secretase complex, Myc-tagged Notch was used as a positive control for Presenilin-1 interaction. As shown in Fig. 7A, HA-tagged p190RhoGEF was specifically co-immunoprecipitated by RhoA in the absence of  $\delta$ -catenin (an arrowhead in the upper first panel). However,  $\delta$ -catenin overexpression noticeably decreased the interaction between p190RhoGEF and RhoA (lane 6 versus 7 in upper first panel). Likewise, RhoA overexpression decreased the interaction between p190RhoGEF and  $\delta$ -catenin (lane 4 versus 7 in the upper third panel). In contrast to the published result showing p120<sup>ctn</sup> directly interacts with RhoA, we could not detect such an interaction between  $\delta$ -catenin and RhoA (upper second panel). Overexpressing  $\delta$ -catenin T454A mutant did not alter the interaction between p190RhoGEF and RhoA (Fig. 7B). These results strongly support our hypothesis that the overexpressed  $\delta$ -catenin lowers the active form of RhoA by sequestering p190RhoGEF to prevent it from activating RhoA, rather than by direct interaction between  $\delta$ -catenin and RhoA.

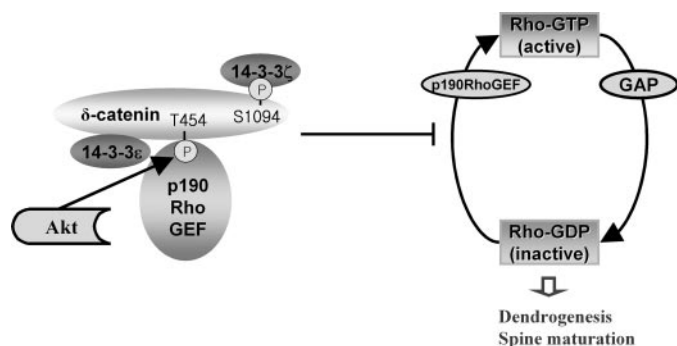
## DISCUSSION

Because dendritic spines are actin-rich protrusions that form the major postsynaptic sites of the excitatory synaptic input, dendrogenesis, and spine morphogenesis are essential for synaptic plasticity and cognitive function. Moreover, spine abnormalities have been shown to be associated with many neurolog-





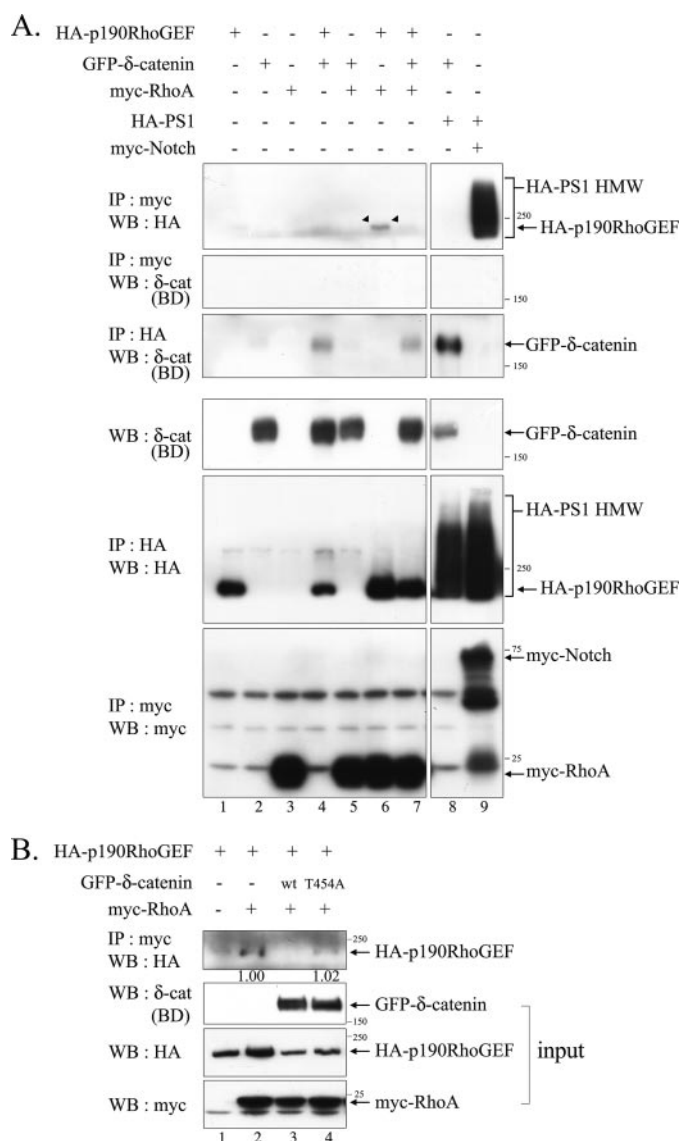
**FIGURE 5. Effects of wild type and mutant  $\delta$ -catenin on the dendrogenesis and spine formation in primary hippocampal neurons.** The cultured hippocampal neurons at 16 DIV were transfected with the full-length GFP- $\delta$ -catenin or various mutants, fixed and stained with GFP antibody. The high magnification images are inverted for clarity. The average number and length of mature spines, either a cotyloid appearance or flat-apex mushroom appearance, were analyzed, and filopodia-shaped spines are excluded from analysis (\*,  $p < 0.01$  by ANOVA and Tukey's HSD post hoc test). The number of dendritic branches that intersects a sphere, 100  $\mu\text{m}$  in radius, centered at the soma was added for plotting and statistical analysis using ANOVA and a Tukey's HSD post hoc test. The graphs show the average number of intersections between the dendritic branches and a sphere (\*,  $p < 0.01$ ). The study has been repeated with three different cultures (3 different litters from 3 different pregnant rats) and from each litter, 3–4 coverslips (~10 neurons per coverslip) have been included in this analysis. Scale bars, 20  $\mu\text{m}$  for low magnification, 3  $\mu\text{m}$  for high magnification, and 1  $\mu\text{m}$  for high magnification of the dendritic spines.



**FIGURE 6. A proposed model of  $\delta$ -catenin-induced morphogenesis.**  $\delta$ -Catenin binds to p190RhoGEF through Akt-mediated Thr-454 phosphorylation, sequesters p190RhoGEF to prevent it from activating RhoA, and thereby reduces the local endogenous RhoA activity. The reduced local GTP-RhoA affects its downstream effectors, which, along with other  $\delta$ -catenin-induced changes, regulates the dendrogenesis and mature spine formation. The interaction between  $\delta$ -catenin with the 14-3-3 isoforms occurs through a different binding domain. The C-terminal region of  $\delta$ -catenin is important for binding to 14-3-3.

ical disorders including most types of mental retardation (19, 43–45). Even though we previously reported that a hippocampal neuron overexpressing  $\delta$ -catenin demonstrates an elaborate arborization of dendrites, swelling, and enhanced dendritic spine maturation (15), a complete picture of how  $\delta$ -catenin affects dendrogenesis and spine morphogenesis would be very important for understanding its role in synaptic plasticity and cognitive function. In this study, we found that  $\delta$ -catenin interacts with p190RhoGEF and significantly lowered the level of GTP-RhoA, and that Akt1 mediates the interaction between  $\delta$ -catenin and p190RhoGEF through Thr-454 phosphorylation of  $\delta$ -catenin. The formation of  $\delta$ -catenin-induced dendrite-like process formation in NIH 3T3 fibroblasts was totally abolished by substitution of Thr-454 residue to Ala, which is a defective form in binding p190RhoGEF. Furthermore,  $\delta$ -catenin T454A significantly reduced the length and number of mature mushroom-shaped spines in primary hippocampal neurons. Overall, these results suggest that the interaction between  $\delta$ -catenin and p190RhoGEF can act as a key modulator in regulating  $\delta$ -catenin-induced dendrogenesis and spine formation.

Our protein motif-scan analysis demonstrated 3 putative sites (Ser-282, Thr-454, Ser-1094) in mouse  $\delta$ -catenin as a putative target phosphorylation sites for Akt and/or 14-3-3 binding sites. The precise mapping for the phosphorylation site by Akt, and subsequently, a binding site for p190RhoGEF and/or 14-3-3, is of great interest because the three molecules,  $\delta$ -catenin, p190RhoGEF, and 14-3-3, may interact with each other independently. For example, a recent report demonstrated that mouse  $\delta$ -catenin interacts with 14-3-3 through Ser-1094 in a phosphorylation-dependent manner (30). 14-3-3 also binds to p190RhoGEF through a unique phosphorylation-independent binding site (I(1370)QAIQNL) in p190RhoGEF (29). Hence, this study examined the possibility that  $\delta$ -catenin may bind to p190RhoGEF via 14-3-3 through Ser-1094, forming a trimeric complex. The  $\delta$ -catenin T454A mutant, unable to bind p190RhoGEF, still interacts with both 14-3-3 isoforms,  $\epsilon$  and  $\zeta$ , suggesting that p190RhoGEF binds directly to the domains containing Thr-454 in  $\delta$ -catenin in a Akt-phosphorylation-dependent manner rather than by forming a trimeric complex. As



**FIGURE 7.  $\delta$ -Catenin overexpression decreases the interaction between p190RhoGEF and RhoA.** A, Bosc23 cells were transfected with GFP- $\delta$ -catenin, HA-p190RhoGEF, Myc-RhoA, HA-PS1 wt, and Myc- $\Delta$ EN1 in a various combination as indicated in the upper side of the figure. The binding between p190RhoGEF and RhoA was examined by immunoprecipitation with the anti-Myc antibody, and Western blotting was performed with anti-HA antibody (upper first panel). The binding between RhoA and  $\delta$ -catenin was examined by immunoprecipitation with the anti-Myc antibody, and Western blotting was performed with anti- $\delta$ -catenin (BD) antibody (upper second panel). The bindings between Presenilin-1 with  $\delta$ -catenin or with Notch were examined for positive control experiments (lanes 8 and 9 in upper first and third panels). Expression of each protein was shown through the bottom three panels. B, Bosc23 cells were transfected with either wild-type or mutant  $\delta$ -catenin together with HA-p190RhoGEF and/or Myc-RhoA. The binding between p190RhoGEF and RhoA was examined by immunoprecipitation with the anti-Myc antibody, and Western blotting was performed with anti-HA antibody (upper first panel). 5% volume of each lysate was subjected to Western blot analysis to show the input level of each protein. The relative intensity of the immunoprecipitated HA-p190RhoGEF band in lane 2 versus 4 was determined by normalization against the input band of HA-p190RhoGEF. 3  $\mu$ g of each plasmid was used to transfect cells plated in 100-mm dish at 50% confluency.

shown in Fig. 7, the overexpressed  $\delta$ -catenin noticeably decreased the interaction between p190RhoGEF and RhoA, supporting the direct interaction between  $\delta$ -catenin and p190RhoGEF as proposed in our model in Fig. 6. Similar to the results of Mackie and Aitken (30),  $\delta$ -catenin  $\Delta$ C207 (Ser-1094



deleted) showed very little 14-3-3 $\zeta$  binding. However, we did observe a faint band, possibly because of formation of heterodimers with other 14-3-3 isoforms. In contrast,  $\delta$ -catenin  $\Delta$ C207 interacts well with 14-3-3 $\epsilon$ , suggesting that  $\delta$ -catenin can combine with the isoforms of 14-3-3 through different binding domains. Interestingly, p190RhoGEF has been shown to bind to the  $\beta$ ,  $\gamma$ ,  $\epsilon$ , and  $\eta$  isoforms but not to  $\zeta$ , or  $\tau$  isoforms of 14-3-3 (29). Therefore, even though our results suggests p190RhoGEF binds directly to  $\delta$ -catenin in an Akt-dependent manner, the possibility that 14-3-3 $\epsilon$  has favorable effects on the association between p190RhoGEF and  $\delta$ -catenin cannot be totally excluded.

The Rho signaling pathway has attracted considerable attention for several reasons. First, the dendrite-like process formation and cytoskeletal remodeling induced by overexpressed  $\delta$ -catenin will greatly mimic those shown by the use of the C3 toxin from *C. botulinum*, in which ADP-ribosylates and specifically inactivates Rho (46, 47). We previously reported that rhodamine phalloidine staining of NIH 3T3 fibroblasts produced lower levels of stress fiber-associated actin filaments (15), which is indicative of reduced Rho signaling in fibroblasts. Even though Martinez *et al.* (12) reported that  $\delta$ -catenin enhances the effects of Rho inhibition on neurite branching, there is no evidence showing that  $\delta$ -catenin reduces the level of GTP-RhoA in cells. This report provides the first biochemical evidence exhibiting that overexpressed  $\delta$ -catenin indeed reduces the levels of GTP-RhoA but not those of GTP-Cdc42 or -Rac1, along with the mechanism that Akt phosphorylates the Thr-454 residue in  $\delta$ -catenin and enables its association of p190RhoGEF.

In contrast to the few reports on  $\delta$ -catenin, studies on p120<sup>ctn</sup>, a prototype of the p120<sup>ctn</sup> family protein, have revealed several lines of evidence that may be unique to p120<sup>ctn</sup> or related to  $\delta$ -catenin. p120<sup>ctn</sup> has been shown to primarily enhance the development of spine-like protrusions, which were abolished by deletions in the Arm domain (48, 49). p120<sup>ctn</sup> overexpression in CHO cells increased the activity of endogenous Cdc42 (~3.1-fold) and Rac1 (~1.9-fold) activity, but diminished the Rho activity by ~45% through the Rho family exchange factor Vav2, which has activity for RhoA, RhoG, Cdc42, and Rac1 (50). Consistent with this data, a deletion of the p120<sup>ctn</sup> gene in hippocampal pyramidal neurons reduced the GTP-Rac1 level (~40%) and increased the RhoA level (~160%) (51). In contrast, another report demonstrated that p120<sup>ctn</sup>-null epidermis showed markedly higher GTP-RhoA levels but similar GTP-Rac1 and -Cdc42 levels (52). Our results demonstrate that overexpressed  $\delta$ -catenin reduces the level of GTP-RhoA but not those of GTP-Cdc42 or -Rac1. It might be possible because p190RhoGEF is a brain-enriched and RhoA specific GEF (24). In contrast to the published result that p120<sup>ctn</sup> directly interacts with Rho, we could not observe such interaction between  $\delta$ -catenin and RhoA, indicating that the interaction between  $\delta$ -catenin and p190RhoGEF might be a unique mechanism for  $\delta$ -catenin to lower RhoA activity. The mechanism for  $\delta$ -catenin binding of p190RhoGEF and subsequent alteration of RhoA activation could be through a sequestration model as suggested in this report. However, we cannot

yet rule out the possibility that  $\delta$ -catenin directly alters the exchange activity of the p190RhoGEF.

Compared with the explicit mechanism for  $\delta$ -catenin T454A inability to lower active RhoA and modulate dendrogenesis,  $\delta$ -catenin  $\Delta$ C207 interactions are more complex and needs further investigation. As shown in Figs. 1 and 3,  $\delta$ -catenin  $\Delta$ C207 binds to p190RhoGEF better than  $\delta$ -catenin FL wt but displays less capability in lowering RhoA-GTP. This suggests that the C terminus in  $\delta$ -catenin also plays a pivotal role in lowering RhoA activity. We speculate two possibilities based on published results. First, in addition to the association between  $\delta$ -catenin and p190RhoGEF, another protein(s) bound to the C terminus of  $\delta$ -catenin can be important for inactivating p190RhoGEF to a full extent.  $\delta$ -Catenin binds to the last PDZ domain of S-SCAM through its C terminus, which can affect the association between  $\delta$ -catenin and NMDA receptors and other glutamate receptors. There was a recent report showing that GRIP, a direct binding protein of AMPA receptor, also binds to the C-terminal region of  $\delta$ -catenin, whose interaction can affect LTD (Ref. 53, Ochiishi *et al.*, Proceedings of 36<sup>th</sup> Annual Neuroscience Meeting, 2006). The association between Erbin and the cadherin and catenin complex was mediated by an interaction between the C-terminal binding motifs (DSWV-COOH) of  $\delta$ -catenin and ARVCF (54). Martinez *et al.* (12) showed that a deletion of 99 amino acid (aa) residues from the  $\delta$ -catenin COOH-terminal ( $\Delta$ C99), but not a deletion of 205 aa ( $\Delta$ C205), retains its ability to interact with Cortactin. As RhoA combines with the glutamate receptors at the spine plasma membrane and NMDA receptor activation decreases the RhoA activity (55, 56), the C terminus of  $\delta$ -catenin may affect the Rho activity by interacting with one or more of these newly identified proteins. It also would be interesting to examine whether Cortactin or 14-3-3 $\zeta$  mediates  $\delta$ -catenin-dependent RhoA inactivation and dendritic spine morphogenesis. Alternatively, the C terminus in  $\delta$ -catenin may be important by being specifically localized to a certain membrane junction. Our results showed that  $\delta$ -catenin  $\Delta$ C207, which can interact with p190RhoGEF but possess a relatively low level of GTP-RhoA, showed total abolishment of dendrite-like processes in NIH 3T3 fibroblasts and a significantly lower number of dendrites in cultured hippocampal neurons, highlighting the important role of the C terminus of  $\delta$ -catenin plus its intact Thr-454 residue in regulating dendrite morphogenesis. Compared with the significantly diminished dendrite formation,  $\delta$ -catenin  $\Delta$ C207 has less affect on the formation of mature mushroom-shaped spine in terms of the number and length than  $\delta$ -catenin T454A. This indicates that  $\delta$ -catenin contributes to dendrogenesis and spine formation through some overlapping and unique signaling pathways for each process. The different contribution of the C terminus in  $\delta$ -catenin in dendrogenesis and spine formation will require future study to determine its precise mechanism.

Even though the potential roles of dendrogenesis by  $\delta$ -catenin (12, 15) and the promotion of mature spines in cultured hippocampal neurons (15, 16) have been described, the effects of  $\delta$ -catenin on the number, shape, and length of spines are not completely understood. Interestingly, Elia *et al.* (51) recently reported a role for p120<sup>ctn</sup> in regulating spine and synapse formation. Consistent with our previous description, cultured hip-

pocampal neurons expressing full-length  $\delta$ -catenin showed an increased number of mature mushroom-shaped spines compared with the GFP control, whereas  $\delta$ -catenin T454A showed noticeable increases in the number of long, thin filopodia-shaped spines with a concomitant reduction in the number of mature mushroom-shaped spines.  $\delta$ -Catenin  $\Delta$ C207 showed intermediate changes in the number and length of mature spines between  $\delta$ -catenin full-length and T454A, which strongly suggests that the C terminus plus the intact Thr-454 in  $\delta$ -catenin are essential for its full effects on the head shape and length of the spine, as well as its maturation. Interestingly, the morphological changes in the spine induced by  $\delta$ -catenin T454A were similar to those induced by the dominant negative form of N-cadherin,  $\alpha$ -N-catenin, and RhoGEF (24, 57–59). There is accumulated evidence suggesting the role of cadherin complexes and Rho in spine formation (57, 59, 60), whereas the progressive loss of F-actin induced by a treatment with latrunculin B results in a rough spine head with random protrusion (59). Compared with the full-length  $\delta$ -catenin,  $\delta$ -catenin T454A showed differences in the recruitment of Akt and p190RhoGEF, and in the levels of GTP-RhoA, suggesting their important roles in  $\delta$ -catenin-induced spine morphogenesis. The reduced activities of the Rho downstream effectors, ROCK, LIMK, and its substrate ADF/cofilin and Dia, in neurons promote the formation of dendritic spines or branches (61). Furthermore, the expression of RhoA V14, a constitutively active form, decreased the spine density and length dramatically (62), and the inhibition of Rho using C3 toxin in cortical and hippocampal neurons has also been shown to increase the density and length of some mouse cortical and hippocampal pyramidal neurons in organotypic slices (63). Consistently, we showed that  $\delta$ -catenin FL wt, which binds to p190RhoGEF and reduces RhoA activation, significantly increased the number and length of spines while  $\delta$ -catenin T454A, which cannot binds to p190RhoGEF, significantly decreased the number and length of mature spines. Therefore, it is possible to say that  $\delta$ -catenin exerts its effects on spine formation through very limited spatial sites including junctions containing cadherins or NMDA and/or AMPA glutamate receptor, which allows for local reduction of RhoA, restricted actin reorganization, and its effects on other potential binding proteins. As the stimuli that produce various forms of long-term potentiation (LTP) cause a rapid local increase in the extension of filopodia and the formation of new spines at the site of stimulation (19, 64), the strong effects of  $\delta$ -catenin and its mutants on the shape, length, and number of spines suggest its assumptive important role in synaptic plasticity and cognitive function. However, future investigation will be needed to compare the functional roles of  $\delta$ -catenin full-length, and T454A mutant in cognition using transgenic mice.

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## SUPPLEMENTAL DATA

### SUPPLEMENTAL MATERIALS

Constructs – The construction of  $\delta$ -catenin  $\Delta$ N85-325 in pEGFP-C1 has been previously described (15). The expression plasmid for dominant negative (dn) form of p190RhoGEF (Y1003A) was kindly provided by Dr. J.R. Lee (Ewha Womans University, Seoul, Korea) (S1); HA-14-3-3 $\epsilon/\zeta$  by W.-J. Song (Inje University, Daejeon, Korea); Akt wt, kd, myr plasmids by Dr. Jong-Kyung Chung (KAIST, Daejeon, Korea).

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### SUPPLEMENTAL FIGURE LEGENDS

**Fig. S1. The interaction between  $\delta$ -catenin and 14-3-3 $\epsilon$  or 14-3-3 $\zeta$  occurs through a different binding domain.** The MEF cells were transfected with various GFP-tagged  $\delta$ -catenin deletion mutants together with either HA-14-3-3 $\epsilon$  (A) or HA-14-3-3 $\zeta$  (B). To examine the binding of  $\delta$ -catenin and 14-3-3 $\epsilon/\zeta$ , immunoprecipitation was performed with the anti-HA antibody and blotted with the anti-GFP antibody (top panel). The expression of exogenous GFP- $\delta$ -catenin and HA-14-3-3 was confirmed by using anti-GFP (middle panel) or anti-HA antibody (bottom panel), respectively. 3  $\mu$ g of each plasmid was used to transfect cells plated in 100 mm dish at 50% confluency.

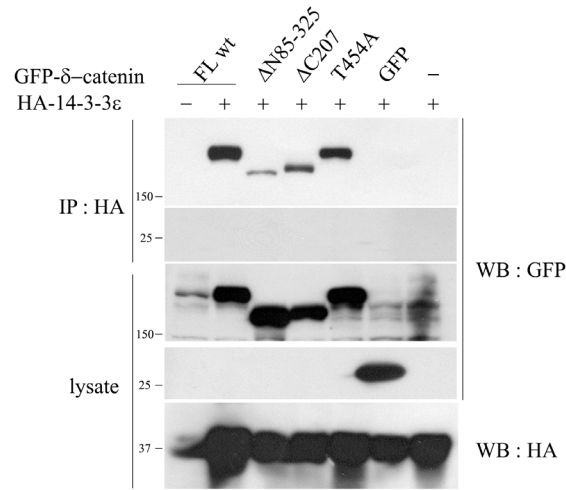
**Fig. S2. The level of  $\delta$ -catenin is increased by both wild type and dominant negative p190RhoGEF in a dose-dependent manner.** The MEF cells were plated in 12 well plate at 50% confluence and transfected with 0.1  $\mu$ g of  $\delta$ -catenin together with either increasing amounts ranging from 0.1  $\mu$ g to 1  $\mu$ g of wild type p190RhoGEF (A) or dominant negative p190RhoGEF (B). The level of  $\delta$ -catenin was examined with anti- $\delta$ -catenin antibody (top panel), and dose-dependent increased expression of HA-

p190RhoGEF was confirmed by using anti-HA antibody (bottom panel). Asterisks indicate non-specific bands. dn, dominant negative.

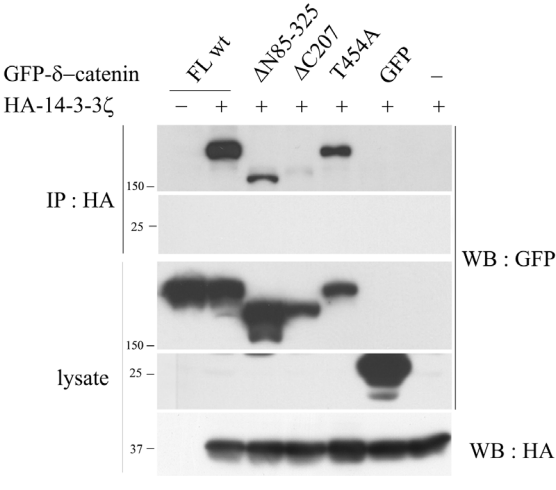
**Fig. S3. Akt1 positively affects the level of  $\delta$ -catenin.** (A) The MEF cells were plated in 12 well plate at 50% confluence and transfected with total 0.2  $\mu$ g of  $\delta$ -catenin together with mock or various Akt1 mutants (wt- wild type, kd- kinase dead, myr- myristoylated form), and the level of  $\delta$ -catenin was determined using anti- $\delta$ -catenin antibody. (B-C) The MEF cells were plated in 12 well plate at 50% confluence and transfected with 0.1  $\mu$ g of  $\delta$ -catenin together with either increasing amounts ranging 0.1, 0.25, and 0.5  $\mu$ g of wild type Akt1 (B) or kinase dead Akt1 mutant (B). The level of  $\delta$ -catenin was determined with anti- $\delta$ -catenin antibody (top panel), and dose-dependent increased expression of HA-Akt1 was confirmed by using anti-HA antibody (bottom panel).

SUPPLEMENTAL FIGURE S1

A.



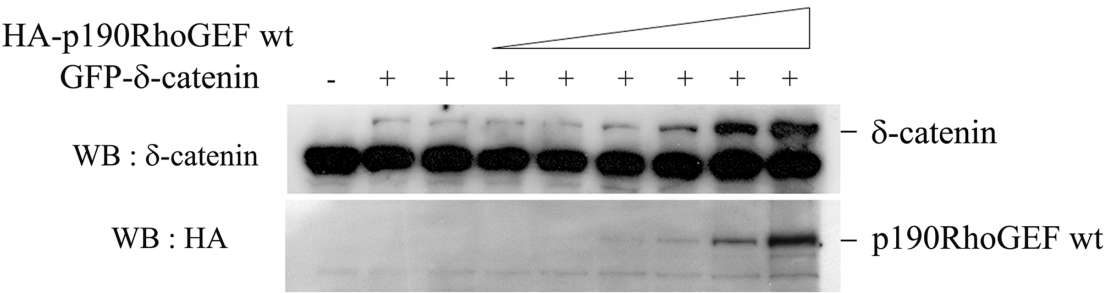
B.



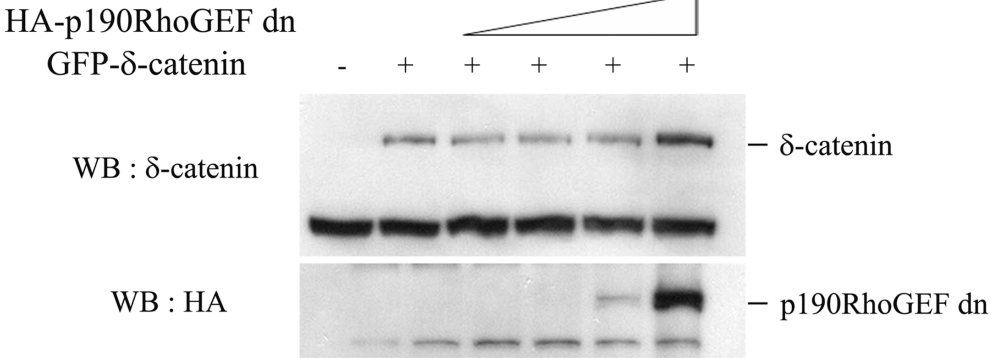


SUPPLEMENTAL FIGURE S2

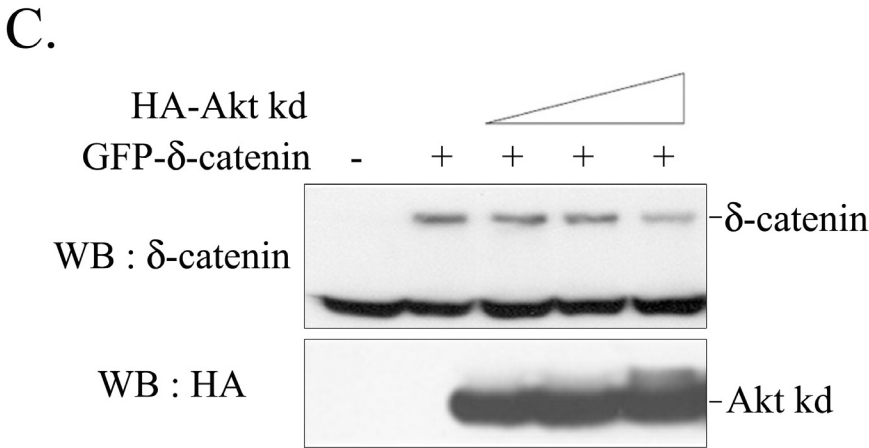
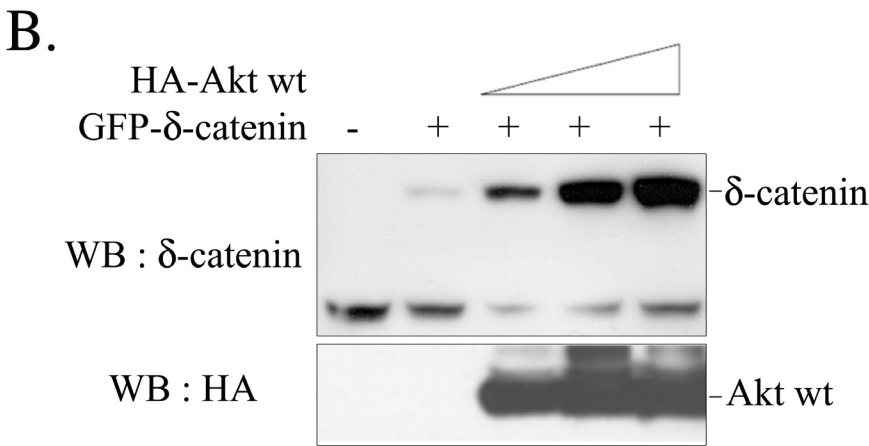
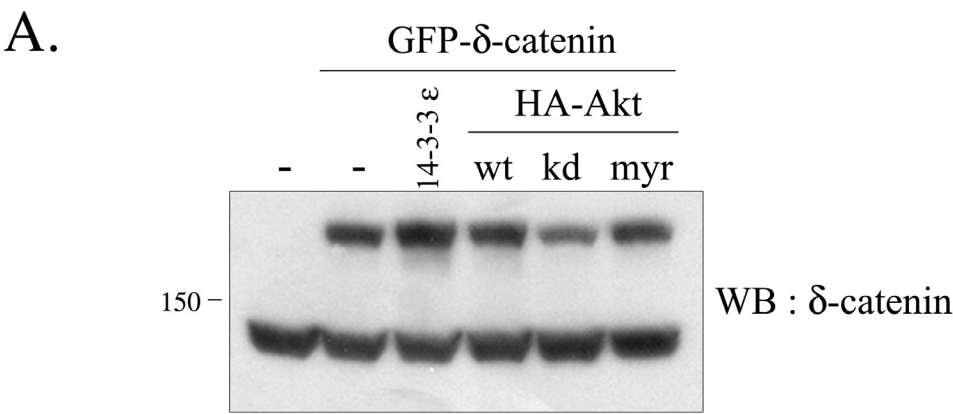
A.



B.



SUPPLEMENTAL FIGURE S3



**$\delta$ -Catenin-induced Dendritic Morphogenesis: AN ESSENTIAL ROLE OF  
p190RhoGEF INTERACTION THROUGH AKT1-MEDIATED  
PHOSPHORYLATION**

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