

Caltubin, a Novel Molluscan Tubulin-Interacting Protein, Promotes Axonal Growth and Attenuates Axonal Degeneration of Rodent Neurons

Nasrin Nejatbakhsh,¹ Cong-Hui Guo,¹ Tom Z. Lu,¹ Lin Pei,^{1,2} August B. Smit,³ Hong-Shuo Sun,^{1,2} Ronald E. van Kesteren,³ and Zhong-Ping Feng¹

Departments of ¹Physiology and ²Surgery, Faculty of Medicine, University of Toronto, Toronto, Ontario M5S 1A8, Canada, and ³Center for Neurogenomics and Cognitive Research, Neuroscience Campus Amsterdam, VU University, 1081 HV Amsterdam, The Netherlands

Axotomized central neurons of most invertebrate species demonstrate a strong regenerative capacity, and as such may provide valuable molecular insights and new tools to promote axonal regeneration in injured mammalian neurons. In this study, we identified a novel molluscan protein, caltubin, ubiquitously expressed in central neurons of *Lymnaea stagnalis* and locally synthesized in regenerating neurites. Reduction of caltubin levels by gene silencing inhibits the outgrowth and regenerative ability of adult *Lymnaea* neurons and decreases local α - and β -tubulin levels in neurites. Caltubin binds to α - and/or β -tubulin in both *Lymnaea* and rodent neurons. Expression of caltubin in PC12 cells and mouse cortical neurons promotes NGF-induced axonal outgrowth and attenuates axonal retraction after injury. This is the first study illustrating that a xenoprotein can enhance outgrowth and prevent degeneration of injured mammalian neurons. These results may open up new avenues in molecular repair strategies through the insertion of molecular components of invertebrate regenerative pathways into mammalian neurons.

Introduction

Whereas adult mammalian peripheral neurons retain the capacity to regenerate injured axons, central mammalian neurons are severely limited in their regenerative growth capacity. The inability of central neurons to regenerate injured axons is due to both extrinsic (Richardson et al., 1980) and intrinsic factors (Sun and He, 2010). Extracellular inhibitory molecules [e.g., Nogo, myelin associated glycoprotein, and OMgp (oligodendrocyte myelin glycoprotein)] limit regeneration in the mature CNS (Mukhopadhyay et al., 1994; Goldberg and Barres, 2000; Wang et al., 2002). However, genetic deletion of these inhibitory molecules is insufficient for successful axonal regeneration (Lee et al., 2010). Manipulation of intrinsic neuronal processes can overcome these extrinsic inhibitory signals and promote axon regeneration both *in vitro* (Qiu et al., 2002; MacGillivray et al., 2009) and *in vivo* (Gao et al., 2004; Steward et al., 2008). Thus, understanding neuron-intrinsic mechanisms of axon regeneration is pivotal in the devel-

opment of treatments for traumatic brain and spinal cord injuries, and neurodegenerative disorders.

In contrast to mature mammalian neurons, adult central neurons of some lower vertebrates and most invertebrates spontaneously regenerate following axonal injury (Ferretti et al., 2003; Mladinic et al., 2009). For instance, adult neurons of the freshwater snail *Lymnaea stagnalis* have the capacity to regenerate their injured axons and reform cell-type specific synapses, both *in vitro* (Syed et al., 1990; Feng et al., 2000; Spencer et al., 2000) and *in vivo* (Koert et al., 2001; Lee and Syed, 2004). A promising avenue to promote axon regeneration in the mammalian CNS may, therefore, be to identify unique proteins that mediate the intrinsic regenerative response in lower organisms and to test the growth-promoting properties of these proteins in the mammalian neurons, where they are not normally expressed. Invertebrate neurons and peripheral mammalian neurons share the capacity to locally synthesize proteins in axons and growth cones, and the identification of locally synthesized proteins has provided valuable insights into mechanisms of axon outgrowth and regeneration (Twiss and van Minnen, 2006). Approximately 100–200 local transcripts have been reported in isolated neurites of various neuronal preparations [e.g., in *Aplysia* sensory neurons (Moccia et al., 2003; Moroz et al., 2006), in squid axoplasm (Capano et al., 1987; Gioio et al., 1994), and in vertebrate axons (Hengst and Jaffrey, 2007; Willis et al., 2007)]. Many of the local transcripts encode cytoskeletal proteins (e.g., microtubule, microfilament, and intermediate filament proteins) (Kaplan et al., 1992; Eng et al., 1999; Vogelaar et al., 2009) or proteins that regulate cytoskeletal dynamics (e.g., Rho and β -thymosin) (Campbell and Holt, 2001; Verma et al., 2005; Wu et al., 2005; van

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Correspondence should be addressed to Dr. Zhong-Ping Feng, 3306 MSB, 1 King's College Circle, Toronto, ON M5S 1A8, Canada. E-mail: zp.feng@utoronto.ca.

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Kesteren et al., 2006), suggesting that regulation of the cytoskeleton through local protein synthesis potentially serves as a conserved mechanism underlying axonal outgrowth and regeneration. Previously, we identified 15 axonally localized mRNAs in *Lymnaea* pedal A (PeA) neurons (van Kesteren et al., 2006). Of these transcripts, one encoded a novel putative calcium-binding protein, caltubin, with no apparent mammalian orthologs or homologs. Here, we present evidence that local synthesis of caltubin promotes axonal regeneration in *Lymnaea* PeA neurons by regulating tubulin level. Expression of caltubin in rat PC12 cells and mouse central neurons attenuates axonal degeneration following injury through a conserved interaction with β -tubulin.

Materials and Methods

Animals. Freshwater snails, *L. stagnalis* (a hermaphrodite), were kept in water at 20°C under a 12 h light/dark cycle and fed green leaf lettuce twice a week (Gardzinski et al., 2007; Hui et al., 2007). Adult snails with shell lengths of 15–20 mm were used. All experiments were performed according to the guidelines of the Animal Care Committee of the University of Toronto.

Primary cell culture and cell isolation. Snails were anesthetized with 10% (v/v) Listerine for 10 min as previously described (Syed et al., 1990; Hui et al., 2007). Central ganglia were dissected in snail saline containing the following (in mM): 51.3 NaCl, 1.7 KCl, 4.1 CaCl₂, 1.5 MgCl₂, pH was adjusted to 7.9 with 1 M HEPES/NaOH, and incubated with 3 mg/ml trypsin (type III; Sigma-Aldrich) for 20 min. Neurons were isolated and placed in poly-L-lysine-coated culture dishes and then maintained in conditioned medium (CM) (Feng et al., 1997) at room temperature up to a week. Anisomycin (A9789; Sigma-Aldrich), a protein synthesis inhibitor that acts by inhibiting peptidyl transferase activity, was added to culture media at various concentrations as previously described (Feng et al., 1997; Roche et al., 2009).

In situ hybridization. Cultured neurons were fixed in 1% paraformaldehyde/1% acetic acid and permeabilized with 0.5% NP-40. Ganglia were fixed in 1% paraformaldehyde/1% acetic acid and embedded in paraffin, and 7 μ m sections were adhered to 0.5% gelatin/0.5% chromalum-coated slides. Digoxigenin-labeled run-off sense and antisense RNA were synthesized from linearized pBluescript plasmids using T3 or T7 RNA polymerase and a digoxigenin-UTP labeling mixture (both from Roche Diagnostics). *In situ* hybridizations were performed as described previously (Smit et al., 1996).

Real-time quantitative PCR. PeA neurons were cultured (8–10 cells per dish) for 2 d. RNA was isolated from the transected neurites and from the corresponding somata separately. After removal of DNA by DNase I treatment, the RNA was random primed with 300 pmol of random hexanucleotides and reverse transcribed into cDNA. Quantitative PCR (qPCR) was performed in triplicate on each cDNA sample using an ABI PRISM 7700 Sequence Detection System (Applied Biosystems) with SYBR Green as the reporter dye. All reactions were performed according to the instructions of the manufacturer. The following primers were used: caltubin (forward, 5'-TTGTAACAGAGGCAGAGCTG-3'; reverse, 5'-AGTTTCCCGTCCTTGTTCCG-3'), thymosin (forward, 5'-CTGTGAGCCATCAAGCTGTTG-3'; reverse, 5'-GGTGTACATGGCATCCGATTT-3'), β -tubulin (forward, 5'-AGCCATCCTTCTGGGTA TG-3'; reverse, 5'-AGTTTCCCGTCCTTGTTCCG-3'), β -actin (forward, 5'-AGCCATCCTTCTGGGATATG-3'; reverse, 5'-ATACCTGGGAAC ATGGTGGT-3'), and mitochondrial 16S rRNA (forward, 5'-ACTTGTAGCTGTGCTAAGGTAGCATAA-3'; reverse, 5'-CAGTTCCTCCCTAT TAATCCGTTTCAT-3'). Overall cDNA expression levels per sample were normalized to the expression of 16S rRNA. The relative expression level of the genes of interest was calculated by $\Delta\Delta C_T$ methods.

Double-stranded siRNA production and delivery. Caltubin-specific 27-mer siRNAs were designed using SciTools RNAi Design software (Integrated DNA Technologies). TriFECTa control RNA was used as a control in our experiments (Integrated DNA Technologies). The siRNA sequences are as follows: caltubin siRNA1 (LcA1), 5'-GAAACUUUCACU UGAUGAAUUCAG-3'; caltubin siRNA3 (LcA3), 5'-CCACGUUCGA CGAGGUCAAGAATA-3'; caltubin siRNA6 (LcA6), 5'-UGGACACU

GUCCAGGGUUAUUACAT-3'; caltubin siRNA10 (LcA10), 5'-UUAU UGAUGCCAUUGGUUAGAUA-3'; TriFECTa control, 5'-UCACAA GGGAGAGAAAGAGAGGAAGGA-3'. Unless otherwise specified, the caltubin siRNA used in the experiments was LcA3.

For whole-animal RNA knockdown experiments, snails were anesthetized with 10% (v/v) Listerine, and 2 μ l of 20 μ M control siRNA or caltubin siRNA was injected into the head, above the central ganglia using a microliter syringe (Hamilton Company) (Fei et al., 2007; Hui et al., 2007; Lu and Feng, 2011). Ganglia were removed 48 h after injection, and RNA or protein was extracted. The mRNA and protein levels of caltubin were measured using semiquantitative RT-PCR and Western blotting, respectively, to confirm knockdown. For cultured cells, siRNA (7 nM) was directly applied to culture medium, as described previously (Hui et al., 2007; Lu and Feng, 2011).

Neurite outgrowth assays and neurite injury in *Lymnaea* neurons. PeA neurons were cultured in CM for 15 h to allow initiation of outgrowth, and cells with initial neurite lengths between 40 and 60 μ m were used. Caltubin siRNA (7 nM) or control siRNA (7 nM) was then applied to culture medium (designated as $t = 0$) and neurite outgrowth was monitored for a period of 30 h using an Olympus inverted microscope (CK X41) with an Olympus C5050 digital camera. Images were analyzed using ImageTool 3.1 software. The net changes in the neurite lengths were measured at various time points following siRNA application ($t = 0$). Where there were multiple neurites for a given cell, an average length of all neurites belonging to each cell was calculated and counted as one sample. Neurite length was measured from the edge of the soma to the tip of the growth cone.

For neurite injury, PeA cells were cultured in CM for a period of 20 h to allow for neurite outgrowth, and cells with initial neurite lengths between 70 and 90 μ m were used. The neurites were severed using glass micropipettes and then immediately treated with either caltubin siRNA (7 nM) or control siRNA (7 nM) (designated as $t = 0$). Neurite measurements were performed on the proximal (soma-attached) and distal (free) ends of the transected neurites at various time points, as described above.

Immunocytochemistry and confocal microscopy. Antibodies against caltubin were raised in rat against synthetic peptides corresponding to the N-terminal (MERADEFVRSQHRD) and C-terminal (DG-KLSLDEFKTLTLYSP) regions of the protein. Cultured PeA neurons were fixed, permeabilized, and incubated with specific primary antibodies (NCS-1: 1:300, BIOMOL, catalog #BML-NL3750-0100; β -tubulin: 1:300, Sigma-Aldrich, catalog #T0198; α -tubulin: 1:300, Sigma-Aldrich, catalog #T6074; β -actin: 1:300, Sigma-Aldrich, catalog #A1978; caltubin: 1:200), as described previously (Gardzinski et al., 2007; Hui et al., 2007). Cells were then incubated with their respective secondary antibody (488 goat anti-rat, Millipore Bioscience Research Reagents; Alexa Fluor 568 goat anti-mouse, Invitrogen; Alexa Fluor 633 goat anti-rabbit, Invitrogen) for 2 h at room temperature. When cells were incubated with secondary antibodies only, no signal was observed under these conditions. Unless otherwise specified, immunocytochemistry of PeA neurons was conducted 6–8 h following siRNA treatment.

Confocal images were acquired using a TCS SL laser confocal microscope (Leica confocal software, version 2.5; build 1347; Leica Microsystems), as described previously (Gardzinski et al., 2007; Hui et al., 2007). Each Z-plane was 0.5 μ m. Background fluorescence was determined in three regions not containing cells throughout the entire z-stack. All siRNA-treated cells were imaged using the same magnification and laser settings. To quantify antibody staining, the mean amplitude fluorescence intensities (in arbitrary units) were averaged from sections of the z-stack for each region of interest, and NCS-1 was used as a control against which microfilament intensities were measured. Each bar represents the average protein level from all cells under that condition.

Protein colocalization analysis. The analysis was conducted to quantify distribution of the proteins labeled with different fluorescence markers and to measure the degree of overlap of the fluorescence signals within the same pixel. Colocalization suggests a high probability of two proteins co-occurring in close proximity. Confocal images of cells double-stained with antibodies for proteins of interest were scanned at a resolution of 1024 \times 1024 pixels in the selected regions of interest (ROIs) in the neurites and growth cones of PeA cells, using a 63 \times oil-immersion lens

A

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Neurocalcin Delta (Bt) : ---MCKNSKLRPEVMODLLESTDEPT---EHEIQBMWYKGFDRDPCPSGHISMBBPKKLYGNEFFPYGDASKFAEHVPTFDAN--GDGTIDFR--EPTIAL : 89
Neurocalcin Delta (Hs) : ---MCKNSKLRPEVMODLLESTDEPT---EHEIQBMWYKGFDRDPCPSGHISMBBPKKLYGNEFFPYGDASKFAEHVPTFDAN--GDGTIDFR--EPTIAL : 89
Neurocalcin Delta (Dr) : ---MCKNSKLRPEVTQDLNDTDEPT---EHELLEWYKGFDRDPCPSGALSMDBPKKLYGNEFFPYGDASKFAEHVPTFDAN--GDGTIDFR--EPTIAL : 89
Neurocalcin, isoform B (Dm) : ---MCKNSKLRPEVLEELKQNTBET---DABIQBMWYKGFDRDPCPSGHISMBBPKKLYGNEFFPYGDASKFAEHVPTFDAN--GDGTIDFR--BELCAL : 89
Calmodulin (Ac) : ---MADQITTEEQI---AFPKKABSLSDKDG--DGNITTRBELGTVMRSLGQNPTEABLQDMINEVDAD---GNGTIDFP--BELTMM : 73
Calmodulin, isoform B (Dm) : ---MADQITTEEQI---AFPKKABSLSDKDG--DGNITTRBELGTVMRSLGQNPTEABLQDMINEVDAD---GNGTIDFP--BELTMM : 73
Calmodulin (Hs) : ---MADQITTEEQI---AFPKKABSLSDKDG--DGNITTRBELGTVMRSLGQNPTEABLQDMINEVDAD---GNGTIDFP--BELTMM : 76
Calmodulin L32 (Os) : ---MDAKQSVAAAKPSTAKKTASAEFLRN---GSLNAVRLRRVFDLDRNG--DCEITVDELAALDALGLVADRDGLAATVSAVYVPEGAAQLRFEDFD--ALHRL : 98
Caltubin (Ls) : MERAPEVDRSQHRDASEHIVSRGTRRSANGGVPCVTVVPLTSLKLRERLRLDHDNDGQSFEDVAVYIRREKEDVTDQTVAAFLSRRDSN---GNCALDFVPEVYVHDM : 105
                                     6 1 e e 5 5 G 3 E 6 G gtiDF e 6
    
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Neurocalcin Delta (Bt) : ---SVTSS---GKLEQKLRWAFSMYDDDGNGYISKAEMDEIVQAIYKVMVSVMMPEDEBSTEPEKRTTEKIFROMDNRDCKLSLSEFTIRGAKSDPSIVR : 181
Neurocalcin Delta (Hs) : ---SVTSS---GKLEQKLRWAFSMYDDDGNGYISKAEMDEIVQAIYKVMVSVMMPEDEBSTEPEKRTTEKIFROMDNRDCKLSLSEFTIRGAKSDPSIVR : 181
Neurocalcin Delta (Dr) : ---SVTSS---GRLDQKLRWAFSMYDDDGNGYISKAEMDEIVQAIYKVMVSVMMPEDEBSTEPEKRTTEKIFROMDNRDCKLSLSEFTIRGAKSDPSIVR : 181
Neurocalcin, isoform B (Dm) : ---SVTSS---GKLEQKLRWAFSMYDDDGNGYISKAEMDEIVQAIYKVMVSVMMPEDEBSTEPEKRTTEKIFROMDNRDCKLSLSEFTIRGAKSDPSIVR : 181
Calmodulin (Ac) : ARKMK---DTSSEBETIRBAFRVFDKDGNGFTSAEIRHVMT---NNG--BKLTDEEVDENIIRBADIDCDGVNYSBEFTMMTTSK--- : 149
Calmodulin, isoform B (Dm) : ARKMK---DTSSEBETIRBAFRVFDKDGNGFTSAEIRHVMT---NNG--BKLTDEEVDENIIRBADIDCDGVNYSBEFTMMTTSK--- : 149
Calmodulin (Hs) : ARKMK---DTSSEBETIRBAFRVFDKDGNGFTSAEIRHVMT---NNG--BKLTDEEVDENIIRBADIDCDGVNYSBEFTMMTTSK--- : 152
Calmodulin L32 (Os) : GDALFGSLDGA AAAAGEPGGGCGDEBEMREAPFRVFDKDGNGFTSAEIRHVMT---NNG--BKLTDEEVDENIIRBADIDCDGVNYSBEFTMMTTSK--- : 196
Caltubin (Ls) : AAPDY---TLGSAWNEFLKLDITNDLDFVTPBAEIVFAE---AVMSPEEALDVTVOGYMSADANNDGKLSLSEFTIRGAKSDPSIVR : 179
                                     e2 aF 6 D 1g1g563 aE6 6 6 e r D 1 DG 6 eEF
    
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Neurocalcin Delta (Bt) : LLQCDPSSAGQF : 193
Neurocalcin Delta (Hs) : LLQCDPSSAGQF : 193
Neurocalcin Delta (Dr) : LLQCDPSSAGC- : 192
Neurocalcin, isoform B (Dm) : LLQCDPSSH--- : 190
Calmodulin (Ac) : --- : -
Calmodulin, isoform B (Dm) : --- : -
Calmodulin (Hs) : --- : -
Calmodulin L32 (Os) : --- : -
Caltubin (Ls) : --- : -
    
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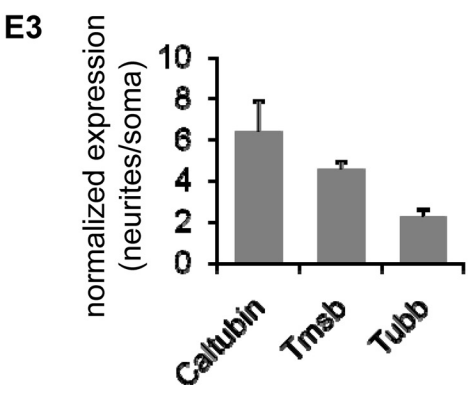
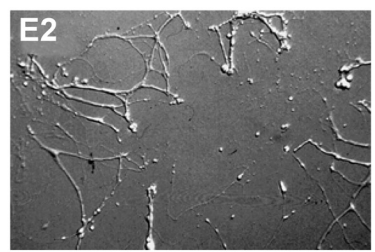
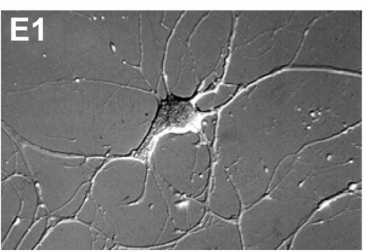
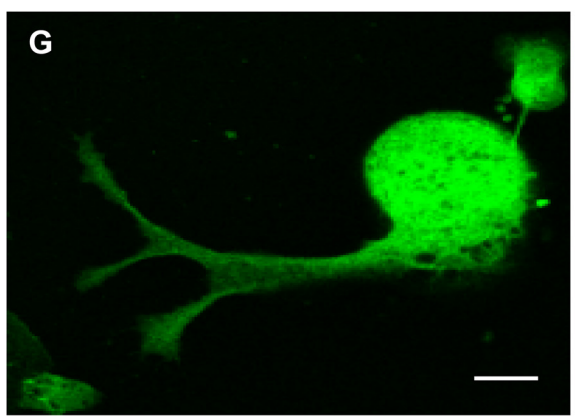
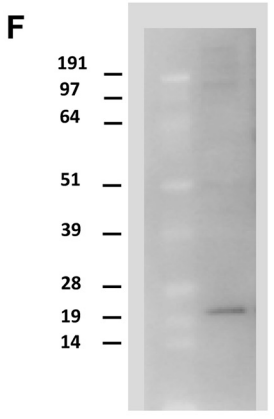
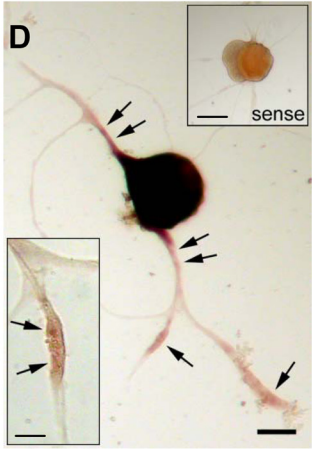
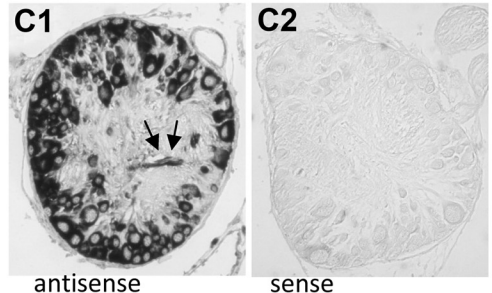
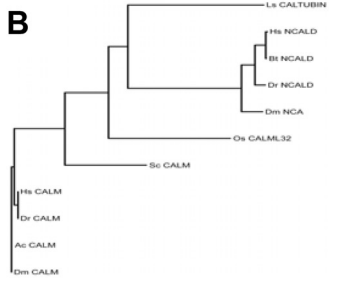


Figure 1. Caltubin is a novel EF-hand-containing protein ubiquitously expressed in CNS neurons of *L. stagnalis*. **A**, The full-length sequence of *caltubin* cDNA predicts a 179 aa protein containing four putative calcium-binding EF-hand motifs (red boxes). GenBank ID numbers are as follows: Neurocalcin Delta: Bt (*Bos taurus*), 5821829; Hs (*Homo sapiens*), 1196122; Dr (*Danio rerio*), 9473279; Dm (*Drosophila melanogaster*) (isoform B), 2724552; Calmodulin: Ac (*Aplysia californica*), 2130764; Dm (isoform B), 7303487; Hs, 825635; Os (*Oryza sativa*) (L32), 7532814; Caltubin: Ls (*Lymanaea stagnalis*). **B**, Phylogenetic tree of the proteins listed in **A**. **In situ** hybridization with a *caltubin* antisense RNA probe (**C1**) revealed presence of *caltubin* mRNA in somata and in neurites (arrows) in the ganglionic neuropil (arrow); or with sense control RNA probes (**C2**). **D**, **In situ** hybridizations of cultured PeA neurons showing differential localization of *caltubin* mRNA in cell soma and neurites labeled by the antisense RNA probe, but not the sense probe (negative control, inset). *Caltubin* mRNA clearly accumulates at specific regions within the neurite, as indicated with arrows. Scale bars: **D**, 20 μ m; right inset, 20 μ m; left inset, 10 μ m. **E**, Quantification of *caltubin* mRNA level in neurites. PeA neurons were cultured for 2 d (**E1**, a representative neuron) and the cell soma was removed (**E2**), allowing mRNA to be isolated from somata and from neurites separately. **E3**, Transcript levels of *caltubin*, β -thymosin (*Tmsb*) and β -tubulin (*Tubb*) (Figure legend continues.)

(Leica Microsystems). Under a zoom of 1, the scan field for each image is $238 \times 238 \mu\text{m}^2$, and thus each pixel is $0.054 \mu\text{m}^2$. Growth cone and neurite were scanned at a zoom of 2. Each fluorescence signal was imaged separately with corresponding channels, and protein colocalization was estimated based on the degree of overlay between the fluorescence intensities, as measured through analysis of each pixel in the ROI, without pixel saturation. For each image, an ROI was selected in the neurite, growth cone, and an area outside the cell to be used as the control. Using the intensity correlation analysis, a Pearson correlation coefficient was calculated using ImageJ software (<http://rsb.info.nih.gov/ij/>), where a coefficient close to 1.0 indicates close colocalization, and 0 indicates low probability for colocalization [also see Li et al. (2004) and Colocalization Module 2007, BioImaging and Optics].

Western blotting. Protein samples were prepared from snail ganglia, mouse brain, or PC12 cells expressing GFP, caltubin-GFP, or caltubin-myc, in RIPA buffer (50 mM Tris-Cl, pH 7.6, 150 mM NaCl, 2 mM EDTA, 1 mM PMSF plus 1% Igepal CA-630, 0.5% sodium deoxycholate, 1% Triton X-100) with a protease inhibitor mixture (5 μl /100 mg tissue; Sigma-Aldrich), as described previously (Fei et al., 2007; Hui et al., 2007). The samples were centrifuged at 4°C at 13,000 rpm for 15 min. Supernatant was extracted and protein concentrations were measured (Bio-Rad). Membranes were incubated with specific primary antibody against caltubin (1:500), β -tubulin (1:1000), α -tubulin (1:6000; Sigma-Aldrich), GFP (1:1000; Ab6673; Abcam), myc-tag (1:500; 05724; Upstate), or β -actin (1:10,000), overnight at 4°C . The membrane was then incubated with appropriate horseradish peroxidase-conjugated secondary antibody accordingly (1:10,000; Jackson ImmunoResearch Laboratories) for 1 h at room temperature. Antibody-labeled protein bands were visualized using enhanced chemiluminescent reagents (PerkinElmer) and analyzed by exposure to film (HyBlot CL).

Coimmunoprecipitation. As described previously (Fei et al., 2007), snail ganglion protein (300 μg) or PC12 cell protein (250 μg) was incubated with Protein A/G Plus-Agarose beads (Santa Cruz Biotechnology) for 30 min on ice. The cell lysates were centrifuged at $10,000 \times g$ for 3 min. The supernatants were collected and then incubated with caltubin primary antibody overnight at 4°C . The protein mix was incubated with Protein A/G Plus-Agarose beads and agitated for 2 h on ice and then centrifuged again at $10,000 \times g$ for 1 min. A sample of supernatant was collected for Western blotting to detect the level of the unbound proteins. The pellet was washed with lysis buffer and centrifuged at $500 \times g$ for 1 min, repeated five times. The pellet was placed in 40 μl of Laemmli buffer (eBioscience; with 50 mM DTT). After vortex, the pellet was boiled for 10 min, and then centrifuged at $10,000 \times g$ for 3 min. The supernatant was collected for Western blotting.

Protein affinity purification (pull-down). For affinity pull-down experiments, GST-caltubin fusion protein was prepared as described previously (Jarvis et al., 2002). Full-length caltubin cDNA was amplified by PCR and cloned into pGEX4T-1. The construct was resequenced to confirm appropriate insertion sites and the absence of spurious PCR generated nucleotide errors. Expression of the transformed plasmid in BL21 (DE3) Bacteria (C25271; New England Biolabs) was induced by IPTG (isopropyl β -D-1-thiogalactopyranoside) (I5502; Sigma-Aldrich). GST-fusion protein was collected from bacterial lysate and purified using glutathione-Sepharose 4B beads as described by the manufacturer (17-0756-01; GE Healthcare). For affinity purification experiments, the solubilized protein extracts (300 μg of protein) were incubated overnight at 4°C with glutathione-Sepharose beads (GE Healthcare) bound to the indicated GST-fusion proteins (50 μg). Beads were washed three times with 600 μl of PBS containing 0.2% Triton X-100, and bound proteins

were eluted with glutathione elution buffer. Eluates were incubated in sample buffer and subjected to 10% SDS-PAGE for Western blot analysis.

PC12 cell culture and transfection. Rat pheochromocytoma PC12 cells were a kind gift from Dr. Shuzo Sugita (University Health Network, University of Toronto, Toronto, Ontario, Canada), who originally obtained the cells from Dr. Thomas Martin (University of Wisconsin, Madison, WI). The cells were maintained on 35 mm poly-L-lysine-coated dishes in a 37°C maintained in a 9% CO_2 incubator in DMEM (HyClone) supplemented with 5% bovine calf serum (HyClone), 5% donor equine (HyClone), 200 U/ml penicillin, 0.2 mg/ml streptomycin, and 0.25 $\mu\text{g}/\text{ml}$ Amphotericin B.

PC12 cells were transfected with caltubin-GFP or GFP plasmid (1–3 $\mu\text{g}/\text{ml}$) using Lipofectamine 2000 (Invitrogen). Six hours following transfection, 50 ng/ml nerve growth factor (NGF) (Invitrogen) was added to the culture medium (DMEM). For neurite outgrowth experiments, cells were maintained in culture for 7 d (DIV7). For the neurite injury experiment, neurite crush was induced on day 5 (DIV5) and cells were monitored for 30 h after injury.

Mouse cortical neuron culture and transfection. All procedures were performed following the animal welfare guidelines at the University of Toronto and were approved by the institutional animal care and use committee. Cortical cultures were prepared from embryonic day 18 mice following the published protocols (Araki et al., 2001; Hares et al., 2011) with modification. Neurons were then dissociated and plated in low density (100,000/ml) on coverslips coated with poly-D-lysine (70–150 kDa; Invitrogen). Cells were maintained in Neurobasal medium with 2% B27 supplement, $1 \times$ Penn/Strep, and 2 mM L-glutamine (all from Invitrogen) at 37°C with 5% CO_2 . Neurons were transfected with caltubin-GFP or GFP plasmid (1–3 $\mu\text{g}/\text{ml}$) using Lipofectamine 2000 (Invitrogen). For neurite outgrowth experiments, cells were transfected on day 1 in culture (DIV1) and maintained for 3 d. For the neurite injury experiment, cells were transfected on DIV3 and neurite were injured on DIV5.

Statistics. Data are presented as the mean \pm SEM. Statistical analysis was performed using SigmaStat 3.0 (Jandel Scientific). The significance of the differences among mean values for multiple experimental groups was tested using one-way ANOVA (Holm–Sidak *post hoc* comparison). Differences were considered significant when $p < 0.05$.

Results

Caltubin transcript and protein are expressed in neurites and growth cones of cultured PeA neurons

We first cloned the full-length cDNA of caltubin (Fig. 1A). Sequencing of a full-length caltubin cDNA predicted a 19 kDa protein with four putative calcium-binding EF-hand motifs (Fig. 1A). Sequence alignment with other calcium binding proteins that show highest sequence similarity with caltubin (i.e., the calmodulins and calcineurins) reveals very low overall sequence identity, implicating caltubin as a novel type of putative EF-hand calcium-binding protein (Fig. 1A,B).

To study the distribution of caltubin transcript in *Lymnaea* neurons, we performed *in situ* hybridization with a caltubin antisense RNA probe, both on pedal ganglia sections and on cultured PeA neurons. In ganglia sections, abundant expression of caltubin mRNA was detected by the caltubin antisense probe in neuronal cell bodies (dark regions) as well as in neurites in the neuropil (arrows) (Fig. 1C1), whereas the sense control RNA probe showed negative in these experiments (Fig. 1C2), indicating the specificity of the RNA probe. In cultured PeA neurons, a strong presence of caltubin mRNA was detected in neuronal somata, and in localized regions of neurites and growth cones by caltubin antisense probe (Fig. 1D), but not by caltubin sense probe (Fig. 1D, right inset). The high-power image (Fig. 1D, left inset) showed that caltubin transcript was unevenly distributed in neurites and accumulated in specific regions (arrows). Real-time

(Figure legend continued.) measured by qPCR analysis are higher in neurites than somata. All mRNA levels were first normalized against 16S rRNA levels, and neuritic mRNA levels were then divided by somatic mRNA levels. The data are presented as mean \pm SEM (n). * $p < 0.05$. **F**, Western blot of total ganglionic protein lysate showing that the N-terminal caltubin antibody recognizes a single protein with a molecular weight of 20 kDa corresponding to the predicted molecular weight of caltubin. **G**, A representative confocal image of a cultured PeA cell labeled with the N-terminal caltubin antibody. Scale bar, 20 μm .

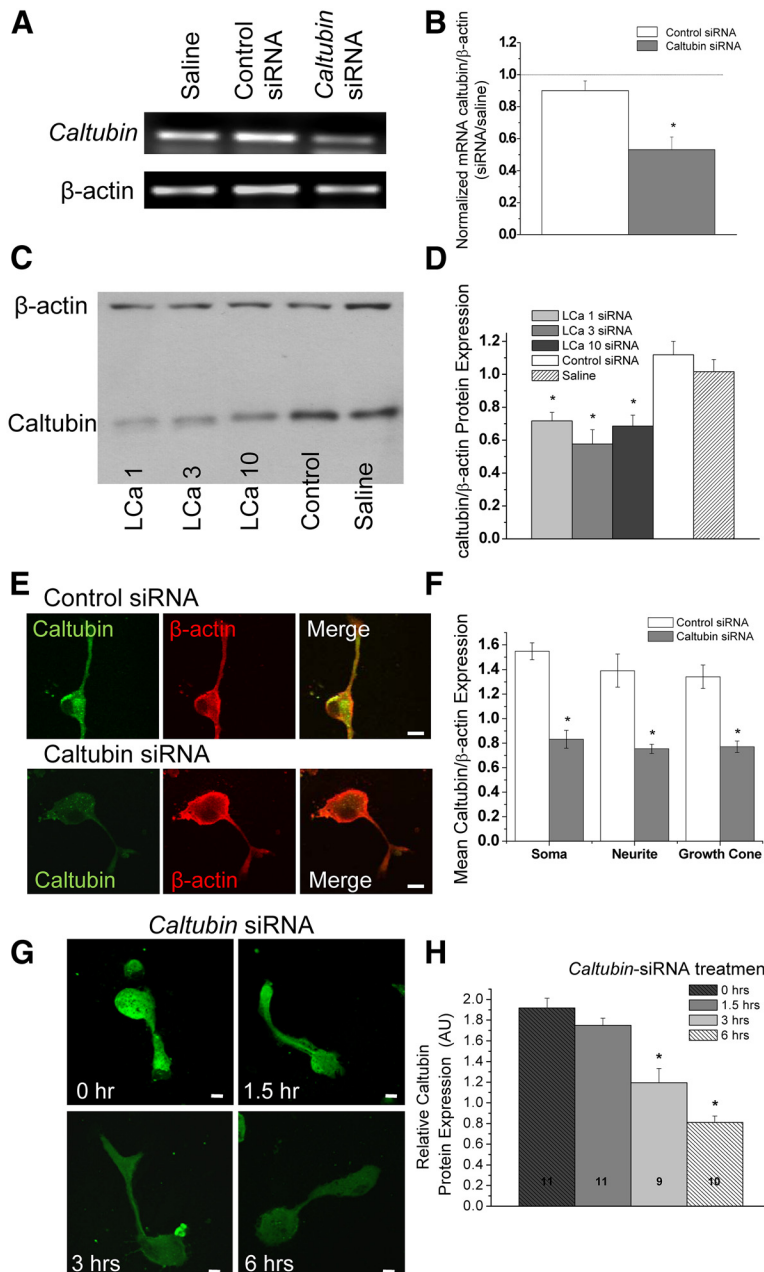


Figure 2. *Caltubin* siRNA treatment significantly reduces expression of both caltubin transcript and protein *in vivo* and *in vitro*. Relative expression levels of ganglial mRNA (**A, B**) or protein (**C, D**) of caltubin against β -actin were measured 24 h following injection of 2 μ l of solution containing saline, control siRNA, or *caltubin* siRNA (LCa1, LCa3, or LCa10) into the snail central ganglia. **A, C**, Representative gel image and Western blot. **B, D**, Quantification of *caltubin* mRNA and protein levels. *Caltubin* siRNA treatment, but not control siRNA treatment, significantly reduced the expression of *caltubin* mRNA ($41 \pm 1.3\%$; $n = 3$) and protein ($54 \pm 1.6\%$; $n = 3$). **E**, Representative confocal images of caltubin and β -actin double-labeled PeA cells cultured in the presence of *caltubin* or control siRNA. **F**, Summary of caltubin to β -actin ratios in soma, neurites, and growth cones of cultured PeA cells treated with control or *caltubin* siRNA, showing a significant reduction in caltubin levels specifically in *caltubin* siRNA-treated cells. Soma: Control siRNA, $n = 11$; *caltubin* siRNA, $n = 10$. Neurites: Control siRNA, $n = 12$; *caltubin* siRNA, $n = 10$. Growth cones: Control siRNA, $n = 12$; *caltubin* siRNA, $n = 11$. **G, H**, Time course of *caltubin* siRNA treatment and caltubin protein expression. **G**, Representative confocal images of immunocytochemical staining of PeA cells at various time points following *caltubin* siRNA (LCa3) treatment. Scale bar, 20 μ m. **H**, A significant reduction in relative caltubin protein levels was already observed at 3 h [1.19 ± 0.13 ($n = 9$; $p < 0.05$)] and 6 h [0.81 ± 0.06 ($n = 10$; $p < 0.05$)] following siRNA treatment, compared with 0 h [1.92 ± 0.94 ($n = 11$)]. All the data are presented as mean \pm SEM (n). * $p < 0.05$.

qPCR analysis confirmed that *caltubin* mRNA is expressed in isolated neurites in culture (Fig. 1E). After 2 d in culture (Fig. 1E1), the cell bodies were removed (Fig. 1E2) and mRNA was extracted from the cell bodies and from neurites separately. Real-

time qPCR analysis was performed to measure the transcript levels of *caltubin* and two other known neuritic transcripts, β -thymosin and β -tubulin (van Kesteren et al., 2006), as positive controls. All mRNA levels were first normalized to the level of 16S rRNA, and neuritic mRNA levels were then divided by somatic mRNA levels. We found that *caltubin* mRNA level was approximately sixfold higher in the neurites than in the cell somata (Fig. 1E3). The neuritic/somatic expression ratio of *caltubin* mRNA was higher than that of β -thymosin and β -tubulin (the positive controls). These data further confirm that *caltubin* mRNA is indeed present at high levels in neurites. To study neuronal caltubin protein expression, we generated antibodies against the C terminus and N terminus of the protein. Immunoblotting showed that all antibodies recognized a single protein band with an apparent molecular weight of ~ 20 kDa corresponding to the predicted molecular weight of caltubin (Fig. 1F). Confocal immunofluorescence imaging revealed caltubin immunoreactivity in the soma, neurites, and growth cones of cultured PeA neurons (Fig. 1G). The presence of both caltubin transcript and protein in neurites and growth cones of cultured adult PeA neurons raises the possibility that local synthesis of caltubin in these distal regions may play a role in regulating neurite development and/or regeneration *in vitro*.

Expression and local synthesis of caltubin are required for neurite outgrowth and regeneration of cultured adult PeA neurons

We next used an RNA interference approach to study the role of caltubin in cultured PeA neurons. Previous studies demonstrated that short siRNAs can be used for efficient gene silencing in cultured *Lymanaea* neurons (Hui et al., 2007). We first showed that *caltubin*-specific siRNAs effectively knocked down caltubin mRNA (Fig. 2A,B) and protein levels, both in intact ganglia *in vivo* (Fig. 2C,D) and in cultured PeA neurons *in vitro* (Fig. 2E,F) 6 h after siRNA application. Three siRNAs (LCa1, LCa3, and LCa10) targeting three different regions of the *caltubin* gene were tested and all resulted in a significant decrease in protein levels (Fig. 2C,D). Caltubin levels measured in cultured PeA neurons progressively decreased over time in *caltubin* siRNA (LCa3) treatment cells, but not in control siRNA-treated cells (Fig. 2G,H).

Caltubin knockdown reached 30% reduction at 3 h (0 h: 1.92 ± 0.09 , $n = 11$; 1.5 h: 1.74 ± 0.07 , $n = 11$; 3 h: 1.19 ± 0.13 , $n = 9$; $p < 0.05$) and 50% reduction at 6 h (0.81 ± 0.06 ; $n = 10$; $p < 0.05$), which was similar to that observed at 24 h.

To elucidate the function of caltubin in neurite outgrowth, we compared neurite extension of cultured PeA neurons in the presence of either *caltubin* siRNA or control siRNA. PeA neurons were cultured for 15 h to allow initiation of outgrowth, and cells with an initial outgrowth of 40–60 μm were used for further analysis. *Caltubin* siRNA or control siRNA were added in the culture medium at $t = 0$, and neurite outgrowth was monitored for 30 h thereafter. We observed a large difference in the outgrowth between the two groups over this period. Neurites of control siRNA-treated cells grew $268.9 \pm 40 \mu\text{m}$ ($n = 17$) on average, which was significantly longer than neurites of *caltubin* siRNA group ($19 \pm 7 \mu\text{m}$; $n = 26$; $p < 0.05$) (Fig. 3A,B). These results indicate that *caltubin* siRNA treatment severely disrupts neurite extension of cultured adult PeA neurons. These findings led us to hypothesize that caltubin might be involved in neurite regeneration.

To test the potential role of caltubin in neurite regeneration of PeA neurons, we transected neurites of cultured PeA cells, and then measured the growth capacity of both the proximal (soma-attached) neurite and the distal (disconnected) neurite (Fig. 3C) following treatment with either control siRNA or *caltubin* siRNA. Specifically, PeA neurons were maintained in culture for 20 h, and cells containing neurites with lengths of $\sim 70 \mu\text{m}$ were used. Neurites were severed using a glass micropipette and immediately treated with either control siRNA or *caltubin* siRNA. Neurite measurements were performed over a period of 30 h following injury (Fig. 3C). As expected, the growth of injured neurites was severely inhibited by *caltubin* siRNA. For proximal injured neurites, a net elongation of $168 \pm 35 \mu\text{m}$ ($n = 7$) was observed in control siRNA condition; in contrast, a net retraction of $23 \pm 4 \mu\text{m}$ ($n = 6$) was observed under the *caltubin* siRNA condition over the same 30 h period ($p < 0.05$; Fig. 3D). These data indicate that caltubin is required for postinjury regeneration of soma attached PeA neurites in culture. Similar to proximal neurites, the distal neurites of transected PeA neurons treated with control siRNA also showed a net increase in neurite length ($133 \pm 5 \mu\text{m}$; $n = 10$) over a 30 h period, whereas the distal neurites of cells treated with *caltubin* siRNA experienced a net retraction of $73 \pm 17 \mu\text{m}$ ($n = 18$) over the same period ($p < 0.05$; Fig. 3E). It is worth noting that all the transected neurites were treated with siRNA after the injury, and the differences observed in the isolated (distal) neurites between control and *caltubin* siRNA treatments over 30 h were likely a result of locally synthesized proteins, which were suppressed by *caltubin* siRNA. To confirm the specificity of our caltubin knockdown, cells were treated for 30 h with each of three *caltubin* siRNAs targeting different regions of the transcript. We found that all three siRNAs consistently reduced

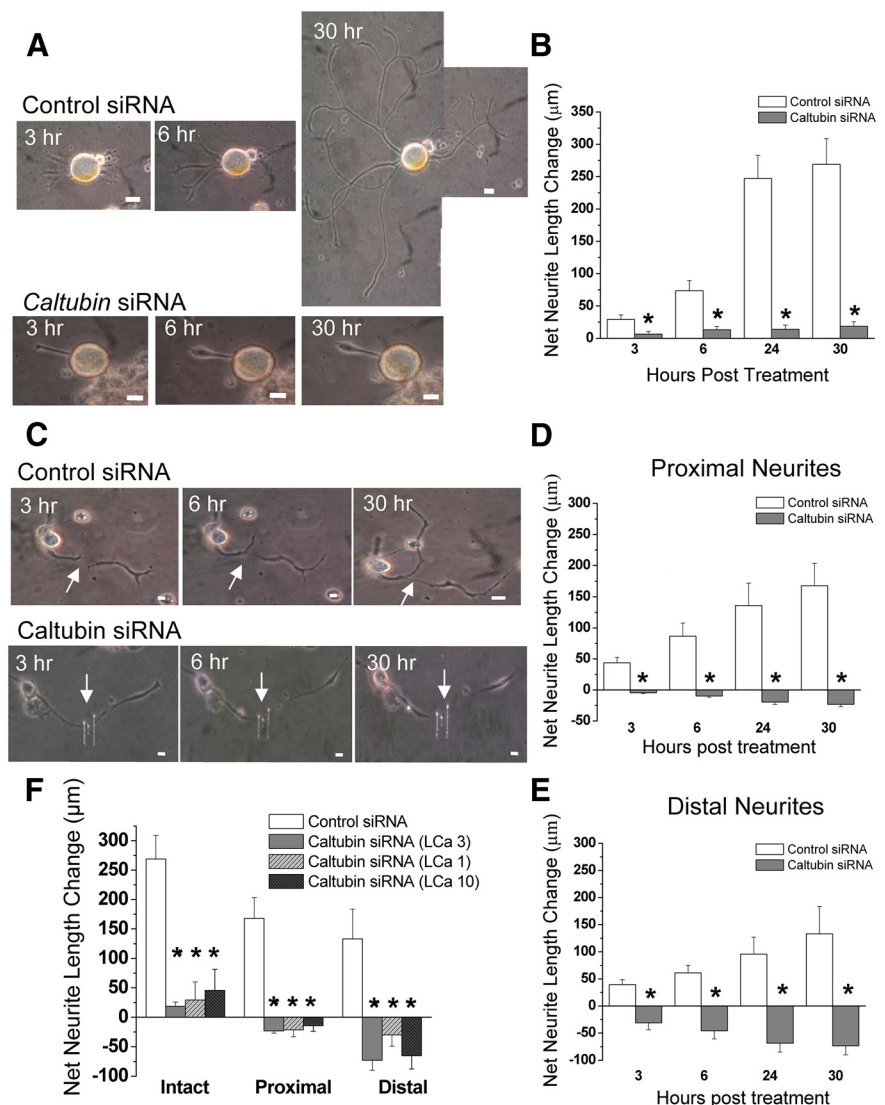


Figure 3. Knockdown of *caltubin* mRNA inhibits elongation of intact neurites and regeneration of injured neurites. **A**, Representative phase-contrast images of a PeA neuron after control siRNA (top) or *caltubin* siRNA application (bottom). **B**, Quantification of the net changes in neurite lengths after siRNA treatment. **C**, Representative images of a transected PeA neuron after control siRNA (top) or *caltubin* siRNA application (bottom). The arrows indicate the site of injury. Quantification of net changes in the proximal (**D**) and distal (**E**) neurite lengths after siRNA treatment. **F**, Three different *caltubin* siRNAs (LcCa3, LcCa1, LcCa10) all block neurite growth of intact, severed proximal, and severed distal neurites. siRNAs were added immediately following neurite transection. The net length of intact (untransected), and the proximal and the distal ends of transected neurites were measured at 30 h after siRNA treatment. * $p < 0.05$. Scale bars: **A**, **C**, 20 μm . All data in **B**, **D**, **E**, and **F** are presented as mean \pm SEM.

the outgrowth ability of the intact neurites and caused substantial retraction in the injured neurites (Fig. 3F). These observations indicate that caltubin is an important protein for normal neurite development, and we therefore decided to address the possible mechanisms by which caltubin might affect neurite outgrowth.

Caltubin regulates β -tubulin but not β -actin protein levels in PeA neurites

Neurite outgrowth and maintenance are highly dependent on the proper assembly of actin and tubulin (Campanot et al., 2003), and can be regulated by calcium-binding proteins (Hui et al., 2007). We investigated the possibility that caltubin affects neurite outgrowth and regeneration by regulating actin or tubulin expression. We conducted immunocytochemical triple labeling of

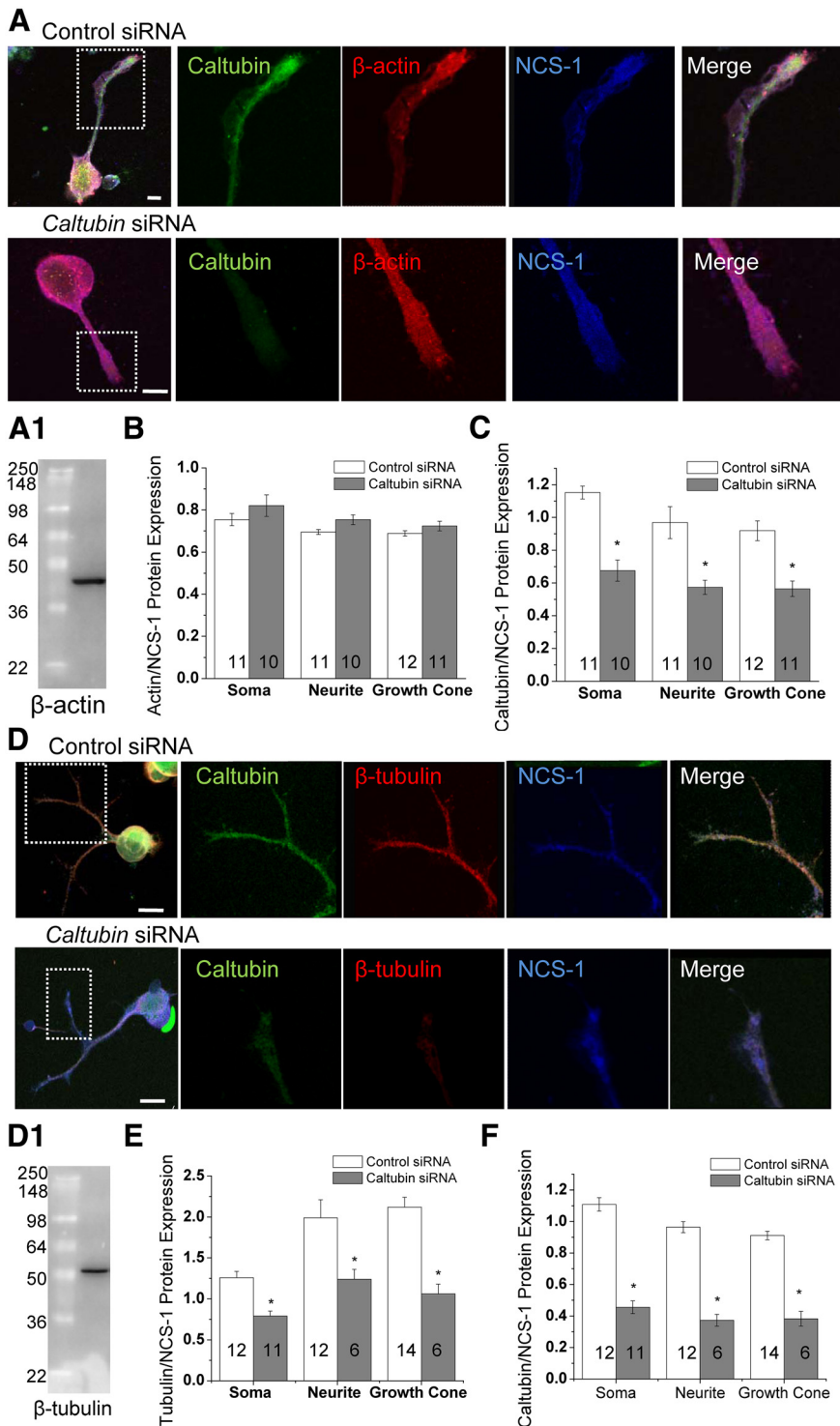


Figure 4. Knockdown of caltubin reduces β -tubulin protein levels, but not β -actin or NCS-1 protein levels. **A**, Representative confocal immunofluorescence images obtained from PeA cells treated with control or *caltubin* siRNA. Caltubin (green), β -actin (red), and NCS-1 (blue) are shown. **A1**, Western blot showing a single protein band labeled by the β -actin antibody used in **A**. Summary of the relative levels of β -actin (**B**) or caltubin (**C**) against NCS-1. **D**, Representatives of triple stainings with caltubin (green), β -tubulin (red), and NCS-1 (blue) in cells treated with control siRNA or *caltubin* siRNA. **D1**, Western blot showing a single protein band labeled by the β -tubulin antibody used in **D**. Summary of the relative protein expression levels of β -tubulin (**E**) or caltubin (**F**) against NCS-1. Scale bar, 20 μ m. The data are presented as mean \pm SEM. * $p < 0.05$.

caltubin, neuronal calcium sensor 1 (NCS-1), and β -actin or β -tubulin in cultured PeA cells treated with either control siRNA or *caltubin* siRNA. NCS-1 was used as a reference protein. We found that the protein expression levels of β -actin and NCS-1 in

the soma, neurites, and growth cones of PeA neurons were not significantly different between control siRNA ($n = 11, 11, 12$)- and *caltubin* siRNA ($n = 10, 10, 11$)-treated cells (Fig. 4A,B), whereas the relative level of caltubin to NCS-1 was significantly reduced (Fig. 4C). These data are consistent with our immunoblot analysis, which showed that global β -actin levels did not vary with *caltubin* siRNA treatment (Fig. 2). In contrast, the relative β -tubulin protein levels in *caltubin* siRNA-treated cells were significantly reduced by $45 \pm 2.7\%$ ($n = 11$; $p < 0.05$) in the soma, $34 \pm 5.8\%$ ($n = 6$; $p < 0.05$) in the neurite, and $30 \pm 9.7\%$ ($n = 6$; $p < 0.05$) in growth cones compared with control siRNA-treated cells ($n = 12, 12, 14$) (Fig. 4D,E), corresponding with a significant decrease in caltubin level (Fig. 4F). Together, these data demonstrate that caltubin has a differential effect on the expression of cytoskeletal proteins. Knockdown of caltubin specifically reduced β -tubulin protein levels, without affecting β -actin protein levels, suggesting that the effect of caltubin on neurite outgrowth and regeneration may be, at least in part, through the regulation of β -tubulin levels.

Neurite outgrowth of PeA cells requires local synthesis of caltubin

Three lines of evidence suggested that caltubin might be synthesized locally in growth cones: (1) *caltubin* transcripts were present in neurites (Fig. 1D,E3), (2) *caltubin* siRNA reduced the caltubin level in growth cones (Fig. 2F), and (3) isolated (distal) neurites treated with *caltubin* siRNA after the injury showed retraction, in contrast to those treated with control siRNA, which showed elongation (Fig. 3E). Because the distal neurites no longer received proteins transported from cell somata, the proteins required for outgrowth of the isolated neurites were likely synthesized locally. *Caltubin* siRNA reduced caltubin level in growth cones (Fig. 2), which may be a result of interruption of the local synthesis of caltubin. To test this hypothesis, we measured the relative levels of caltubin over NCS-1 (as a control) in transected neurites of the PeA neurons either immediately or 3 h following injury (Fig. 5A). The levels of caltubin in both distal (Fig. 5B) and proximal (Fig. 5C) neurites increased significantly within 3 h following injury (distal: 0 h, 0.56 ± 0.05 , $n = 8$; 3 h, 0.71 ± 0.05 , $n = 7$; $p < 0.05$; proximal: 0 h, 0.45 ± 0.03 , $n = 8$; 3 h, 0.72 ± 0.05 , $n = 7$; $p < 0.05$). The increase in caltubin levels in distal neurites supports the hypothesis that the pro-regenerative effects of caltubin are, at least in part, derived through local synthesis of caltubin.

To confirm that transected neurites indeed synthesize caltubin locally, we compared the expression of caltubin, β -tubulin, and NCS-1 in distal regions of transected neurites, in the presence or absence of a protein translation inhibitor, anisomycin (11.3 μ g/ml) (Fig. 5D). We found that both caltubin and β -tubulin protein levels, but not NCS-1 protein levels, were significantly reduced in distal neurites 30 h following anisomycin treatment compared with the control (Fig. 5E). Outgrowth of the intact neurites and the severed neurites following transection was suppressed and a retraction of severed neurites was observed in the presence of anisomycin (Fig. 5F). These observations further support the notion that neurite outgrowth/regeneration in cultured *Lymnaea* PeA neurons requires local translation of caltubin in the distal regions of neurites.

Caltubin colocalizes and interacts with tubulin but not β -actin in *Lymnaea* neurons

To further explore the mechanism of β -tubulin regulation by caltubin, we first tested the degree of spatial colocalization between caltubin and β -tubulin using confocal immunofluorescence intensity correlation analysis. Protein colocalization suggests a high probability of two proteins co-occurring in close proximity. Colocalization was evaluated by measuring the level of overlap between signal intensities of the corresponding fluorescent channels over the selected pixels in randomly selected regions of PeA neurites and growth cones. PeA cells previously treated with control siRNA were double-labeled for caltubin and either β -tubulin, β -actin, or NCS-1. Colocalization of proteins in neurites and/or growth cones was estimated based on the Pearson correlation coefficient: a coefficient (R_r) close to 1 indicates colocalization, while a coefficient close to 0 indicates no colocalization (Li et al., 2004). Our data show that caltubin is colocalized with β -tubulin in neurites and growth cones (neurites: $R_r = 0.67 \pm 0.08$, $n = 6$; growth cones: $R_r = 0.75 \pm 0.10$, $n = 6$; Fig. 6A). In contrast, a much lower correlation was found between caltubin and β -actin ($R_r = 0.10$ – 0.20 ; Fig. 6B) and between caltubin and NCS-1 ($R_r = 0.18$ – 0.30), indicating that there was a low probability for colocalization between caltubin and β -actin or NCS-1. Protein colocalization only implies a high probability of two proteins occurring together in a specific region. Thus, to further determine whether the colocal-

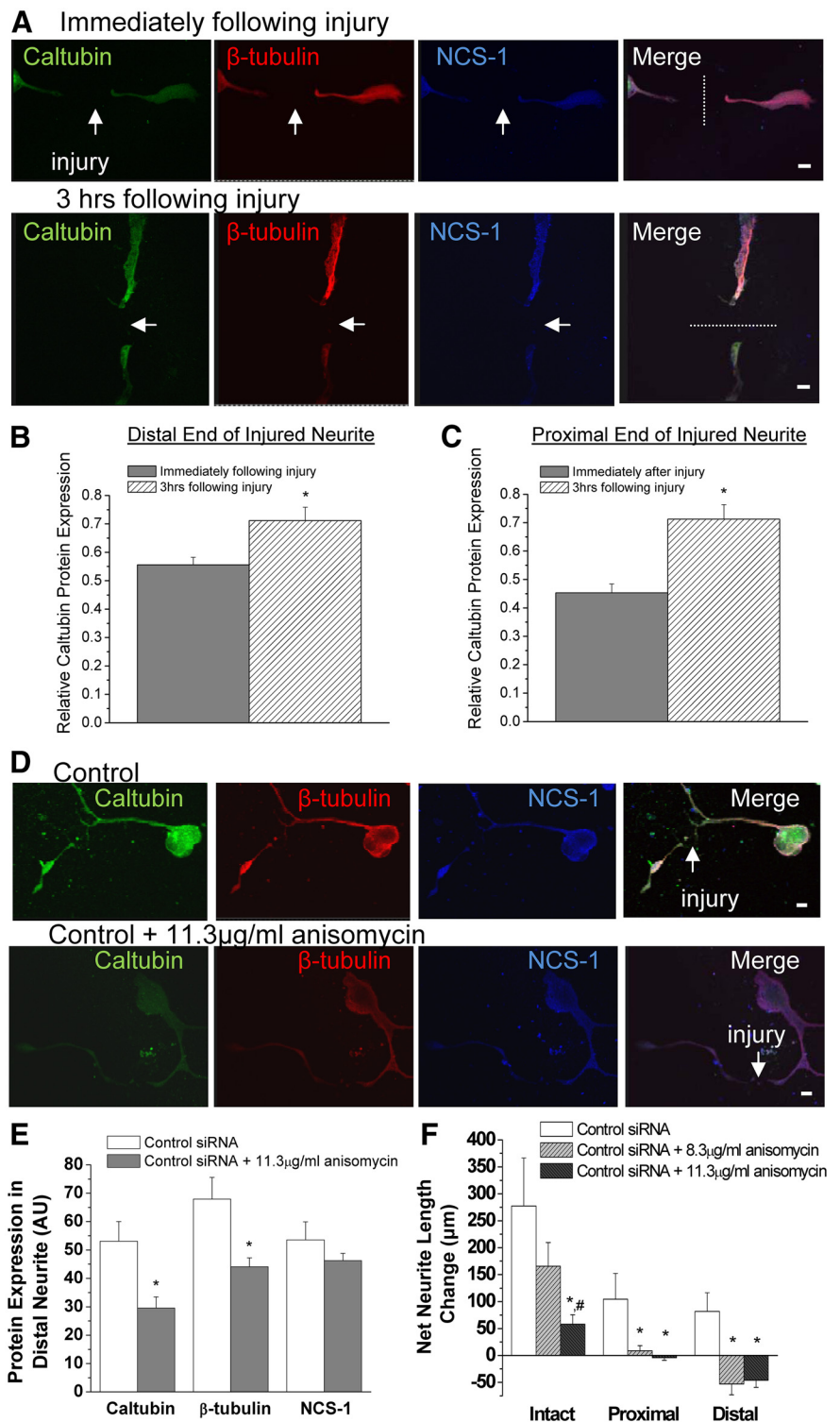


Figure 5. Injury increases caltubin protein levels and blocking protein translation reduces caltubin levels in the distal portion of transected neurites. **A**, Representative confocal immunofluorescence images obtained from transected PeA neuron fixed either immediately following injury (top panel) or following a 3 h delay (bottom panel). Relative levels of caltubin in distal (**B**) and proximal (**C**) portions of neurite measured immediately and 3 h after injury. **D**, Representative confocal immunofluorescence images obtained from injured PeA cells 30 h after treatment with either control siRNA (top panel) or control siRNA plus 11.3 μ g/ml anisomycin (bottom panel). Cells were treated immediately following transection. **E**, Summary of the expression levels of caltubin, β -tubulin, and NCS-1 in the distal neurites with or without anisomycin treatment in presence of control siRNA. **F**, Net changes in neurites following anisomycin treatments. Anisomycin (8.3 or 11.3 μ g/ml) was added in the culture dish immediately after transection. The lengths of the intact neurites and of proximal and distal ends of transected neurites were measured at 30 h after the treatment. Shown are caltubin (green), β -tubulin (red), and NCS-1 (blue). Scale bar, 20 μ m. All the data are presented as mean \pm SEM. * $p < 0.05$.

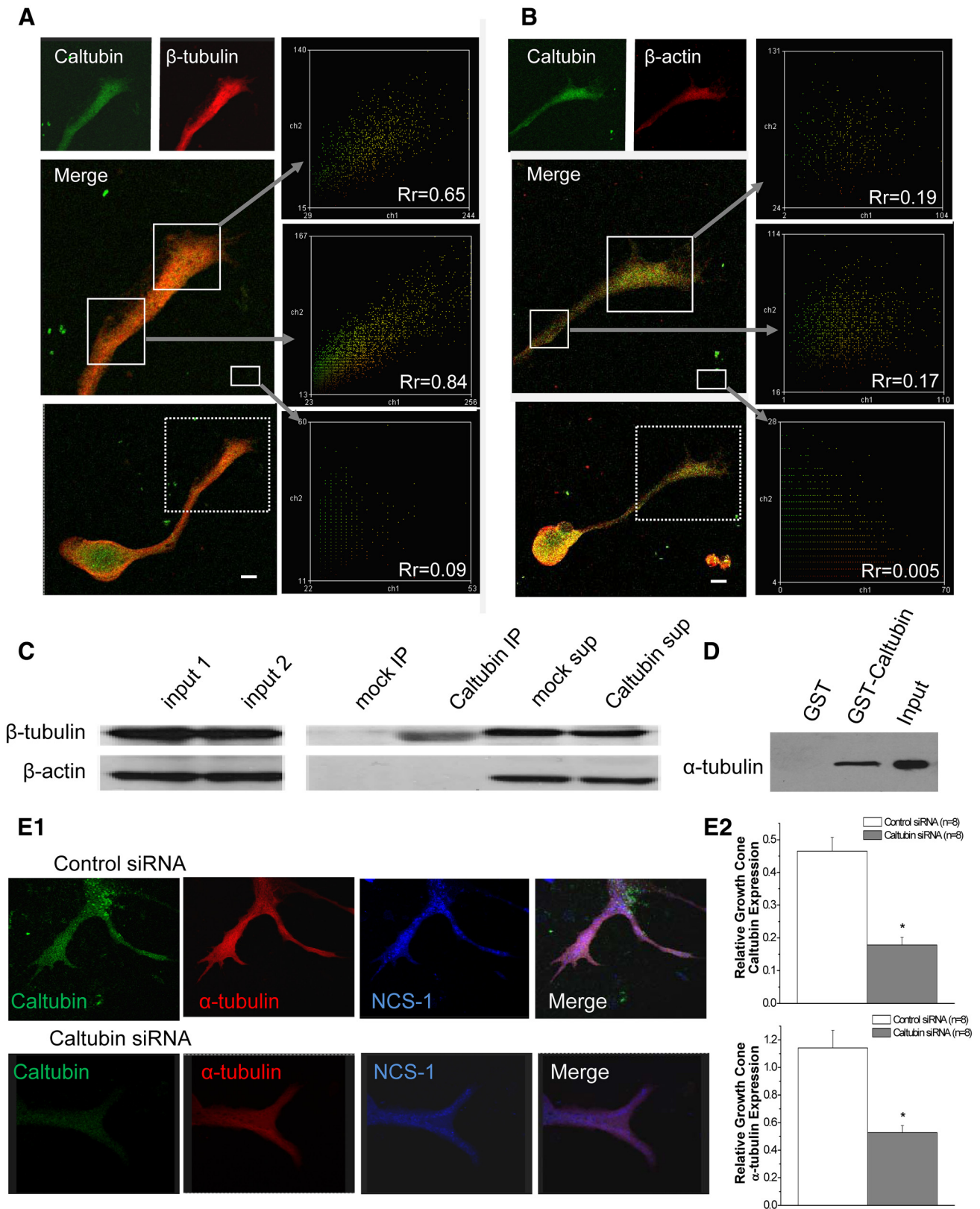


Figure 6. Caltubin is colocalized with and binds to tubulins, but not to β -actin, in PeA neurons. Representative confocal images of double immunocytochemical staining for caltubin and either β -tubulin (**A**) or β -actin (**B**). The correlations between the local staining intensities of the two proteins was measured in neurites, in growth cones, and in the background, and indicated by Pearson coefficient values (R_r). Scale bars, 20 μ m. **C**, Representative Western blots showing that β -tubulin, but not β -actin, coimmunoprecipitates with caltubin. Both β -tubulin and β -actin are detected in input (extracted *Lymnaea* ganglionic protein) as well as in supernatant fractions; however, only β -tubulin is detected in the caltubin IP fraction, indicating specific coimmunoprecipitation between β -tubulin and caltubin. **D**, Representative Western blot of α -tubulin after affinity precipitation revealed a single band in the input and in presence of GST-caltubin, but not with GST alone. **E**, Knockdown of caltubin reduces α -tubulin levels. **E1**, Representative confocal immunofluorescence images obtained from PeA neurons treated with control or caltubin siRNA. Caltubin (green), β -actin (red), and NCS-1 (blue) are shown. Scale bar, 7 μ m. **E2**, Summary of the relative levels of caltubin (top panel) or α -tubulin (bottom panel) against NCS-1, from PeA neurons treated with control siRNA ($n = 8$) or caltubin siRNA ($n = 8$). All the data are presented as mean \pm SEM. * $p < 0.05$.

ized proteins physically interact, we performed a series of immunoprecipitation (IP) or affinity pull-down analyses. We found that caltubin bound to β -tubulin (Fig. 6C), but not to β -actin (Fig. 6C) or the negative control IPs (Fig. 6C). Interestingly, we also observed caltubin binding to α -tubulin (Fig. 6D). To further investigate whether caltubin also regulates α -tubulin levels in growth cones, we used triple immunocytochemical staining (Fig. 6E1). Similar to β -tubulin, relative α -tubulin protein levels in growth cones of *caltubin* siRNA-treated cells were significantly reduced (0.53 ± 0.05 ; $n = 8$) compared with control siRNA-treated cells (1.14 ± 0.13 ; $n = 8$; $p < 0.05$) (Fig. 6E2, top panel) and corresponded with a significant decrease in caltubin levels (caltubin siRNA: 0.18 ± 0.024 , $n = 8$; control siRNA: 0.47 ± 0.042 , $n = 8$; $p < 0.05$) (Fig. 6E2, bottom panel). Together, these results indicate that caltubin colocalizes and interacts with tubulins in *Lymnaea* neurons, and point to a direct mechanism in the regulation of microtubule proteins by caltubin.

Caltubin interacts with β -tubulin, promotes axonal elongation, and attenuates neurite retraction of PC12 cells

Since tubulins are highly conserved (Burns and Surridge, 1990), we asked whether the mechanism of β -tubulin regulation by caltubin might translate to mammalian cells. We first tested the potential binding between caltubin and β -tubulin in rat PC12 neurons by conducting coimmunoprecipitation experiments on cultured PC12 cells, which had been transfected with caltubin-myc, caltubin-eGFP, or eGFP only (each construct at $3 \mu\text{g/ml}$) and allowed a period of 3 d for growth. Our Western blot data show that β -tubulin was present in all three input samples (Fig. 7A1); however, the relative levels of β -tubulin in caltubin-expressing cells over GFP-expressing cells significantly increased (caltubin-GFP: 1.17 ± 0.07 ; caltubin-myc: 1.12 ± 0.04 ; $n = 4$; $p < 0.05$) (Fig. 7A2). In immunoprecipitated samples, β -tubulin was detected in protein complexes obtained from PC12 cells transfected with caltubin-eGFP and caltubin-myc, but not from cells transfected with only eGFP (Fig. 7A1). To confirm these findings, we measured the level of unbound β -tubulin remaining in the supernatant collected following the immunoprecipitation assay. We found that the level of unbound β -tubulin remaining in the supernatant of the eGFP-transfected cells was higher than that in the supernatant of the caltubin-eGFP and caltubin-myc transfected cells, consistent with our IP data. These experiments indicate that, in mammalian cells, the ability of caltubin to bind to β -tubulin is conserved.

Next, we tested whether caltubin can influence axonal elongation in PC12 cells. Cells were transfected with either 1, 2, or 3

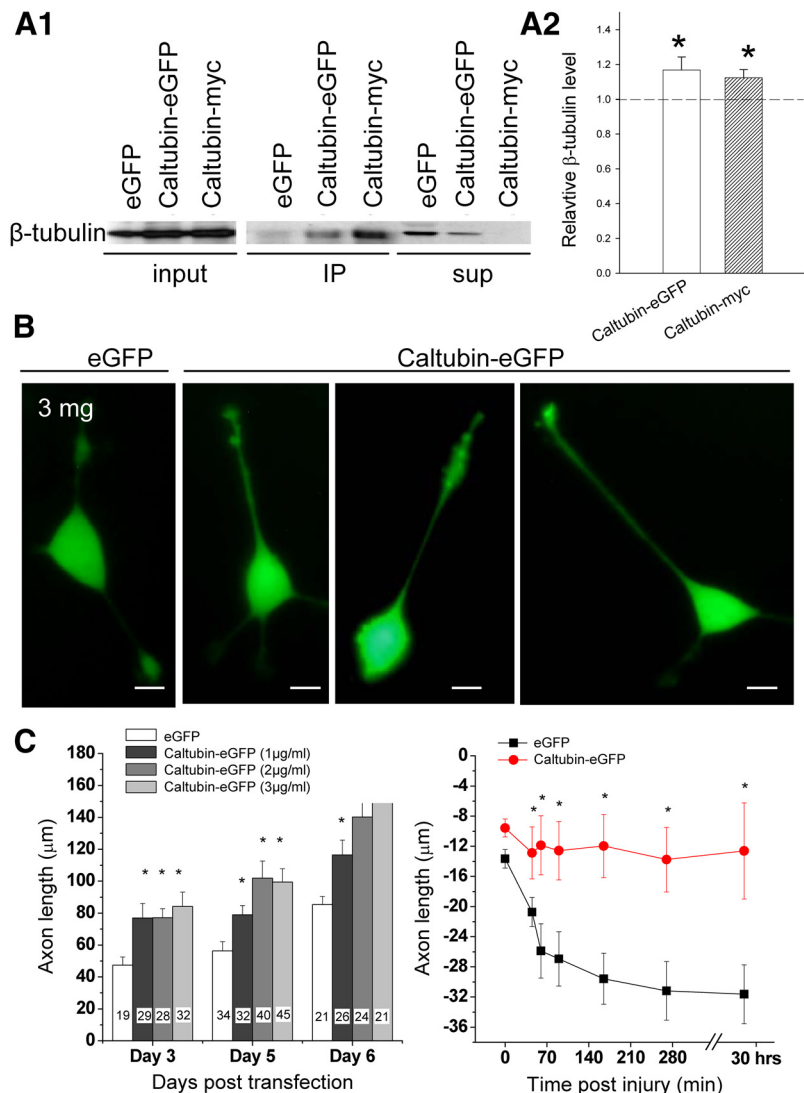


Figure 7. Caltubin expression in rat PC12 cells promotes axonal elongation and attenuates axonal degeneration. **A1**, Representative immunoblots of β -tubulin in cultured PC12 cells transfected with caltubin-GFP, caltubin-myc, or eGFP only (each construct at $3 \mu\text{g/ml}$) and immunoprecipitated (IP) for caltubin. β -Tubulin was detected in all three input samples (lysates), while coimmunoprecipitated β -tubulin was only detected from caltubin-eGFP or caltubin-myc transfected cells. **A2**, Summary of relative levels of β -tubulin in PC12 cells expressing caltubin-GFP or caltubin-myc. The dashed line indicates β -tubulin level in PC12 cells expressing GFP only. **B**, Representative images of PC12 cells expressing caltubin-eGFP or eGFP only. Scale bar, $20 \mu\text{m}$. **C**, Summary of NGF-dependent axonal growth of PC12 cells expressing eGFP, or caltubin-eGFP measured over a period of 6 d after transfection. **D**, Time course of the postinjury changes in axon lengths of PC12 cells expressing eGFP or caltubin-eGFP. Cells were transfected with either caltubin-eGFP ($3 \mu\text{g/ml}$) or eGFP only. Injury was induced by cutting off the growth cone on day 5 after transfection. All the data are presented as mean \pm SEM. * $p < 0.05$.

$\mu\text{g/ml}$ caltubin-eGFP construct or $3 \mu\text{g/ml}$ eGFP control construct (Fig. 7B), and NGF was added on the first day following transfection. Consistent with previous reports, PC12 cells extended processes following NGF exposure (Greene and Tischler, 1976) and were capable of axonal elongation for up to 10 d in culture (Fujii et al., 1982). Overexpression of caltubin-eGFP significantly increased axon extension compared with the eGFP-only controls, in a time-dependent and dose-dependent fashion (Fig. 7C). Specifically, following transfection, the axons of control cells expressing eGFP extended over time in culture (3 d: $47.34 \pm 5.1 \mu\text{m}$, $n = 19$; 5 d: $56.31 \pm 5.9 \mu\text{m}$, $n = 34$; 6 d: $85.4 \pm 5.1 \mu\text{m}$, $n = 21$). The axons of cells treated with caltubin-eGFP ($1 \mu\text{g}$ of construct) extended significantly longer (3 d: $77 \pm 9.1 \mu\text{m}$, $n = 29$; 5 d: $79 \pm 5.8 \mu\text{m}$, $n = 32$; 6 d: $116.4 \pm 9.3 \mu\text{m}$, $n = 26$;

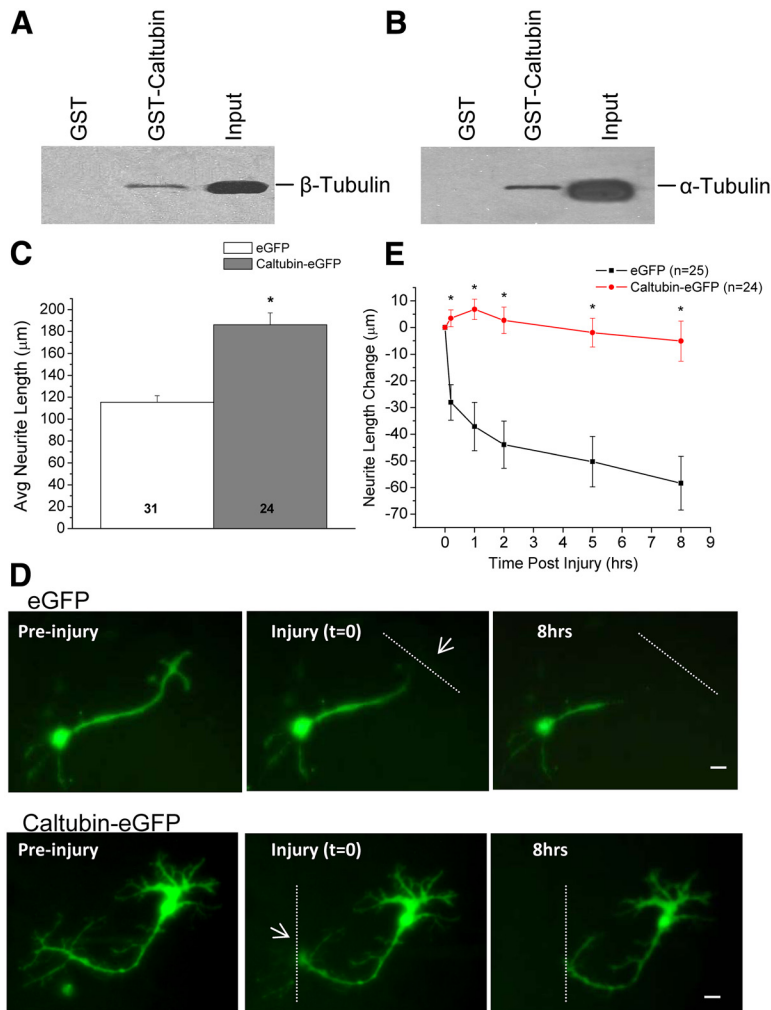


Figure 8. Caltubin expression in mouse primary cortical neurons promotes axonal elongation and attenuates axonal retraction. Representative Western blots of mouse brain β -tubulin (**A**) or α -tubulin (**B**) after affinity precipitation revealed a single band in the presence of GST-caltubin, but not with GST alone. Input (mouse brain tissue-extracted protein) in each experiment serves as the control for indicated proteins. **C**, Summary of NGF-induced axonal growth of mouse primary cortical cells expressing either eGFP, or caltubin-eGFP at DIV3 (transfection at DIV1). **D**, Representative images of cultured cortical neurons expressing eGFP (top panel) or caltubin-eGFP (bottom panel). Neurons were injured at DIV5, 3 d following transfection. Injury was induced by a cut through the growth cone of cultured cells using a glass micropipette, and the dashed line indicates the site and the direction of injury. Scale bar, 20 μ m. **E**, Time course of postinjury changes in axon lengths of cortical neurons expressing eGFP or caltubin-eGFP. All the data are presented as mean \pm SEM. * $p < 0.05$.

$p < 0.05$), compared with the controls. The increase in axonal extension was also observed at higher caltubin-eGFP concentrations (2 and 3 μ g) (Fig. 7*B,C*).

To test whether caltubin can affect axonal degeneration following transection in PC12 neurons, we expressed caltubin-eGFP (3 μ g/ml) or eGFP alone and allowed processes to grow for a period of 5 d. Cells with initial axons of ~ 70 μ m were identified and marked for time-lapse imaging. Axons were injured by cutting off the growth cone using a glass micropipette, and axonal behavior was monitored over various time points following the injury. At 45 min after the procedure, injured axons of cells transfected with eGFP alone showed a significant net retraction of 20.72 ± 1.94 μ m ($n = 19$), whereas axons of cells transfected with caltubin-eGFP showed a net retraction of only 12.89 ± 3.45 μ m ($n = 15$) ($p < 0.05$; Fig. 8*D*). The axons in the eGFP-only group continued to retract over the next 24 h (total net retraction of 31.62 ± 3.9 μ m; $n = 19$), whereas axons expressing caltubin-

eGFP remained stable over the same period of time (total net retraction of 12.62 ± 6.4 μ m; $n = 15$) ($p < 0.05$; Fig. 8*D*). Together, these results demonstrate that caltubin can interact with endogenous mammalian β -tubulin to promote NGF-induced outgrowth and prevent retraction of neuron-like processes following injury in mammalian cells.

Caltubin interacts with tubulin, promotes axonal elongation, and attenuates axonal retraction of cultured mouse cortical neurons

To confirm that caltubin indeed has a conserved role in mammalian neurons, we tested whether caltubin binds to tubulin and promotes neurite outgrowth in mouse cortical neurons. Our affinity pull-down assays showed that GST-caltubin fusion protein, but not GST, precipitated with both β - (Fig. 8*A*) and α -tubulin (Fig. 8*B*), indicating that caltubin binds to mouse microtubule proteins. We then expressed caltubin-eGFP in neonatal cortical neurons on the first day in culture (DIV1) and measured neurite lengths of neurons expressing either eGFP or caltubin-eGFP at DIV3. Neurons expressing caltubin-eGFP had significantly longer average neurite lengths (186 ± 11 μ m; $n = 24$) compared with neurons expressing eGFP only (115 ± 6 , $n = 31$; $p < 0.05$) (Fig. 8*C*), indicating that the snail protein caltubin enhances the growth capability of the mouse neurons. To test whether caltubin also has an effect on neurite regeneration/degeneration, we transfected neurons at DIV3 and cut through the growth cone of one of the neurites at DIV5. We then measured the net changes in neurite length for a period of 8 h following injury (Fig. 8*D,E*). We observed that cells expressing caltubin-eGFP were able to maintain their neurite length following injury over the 8 h period (8 h: $-5.11 \pm$

7.4 μ m, $n = 24$), whereas those expressing eGFP only showed neurite retraction (8 h: -58.38 ± 10.1 μ m, $n = 25$; $p < 0.05$) (Fig. 8*D*). These data confirm our findings in PC12 cells and indicate that caltubin serves to promote neurite outgrowth and attenuate neuronal retraction in mammalian neurons, where it is normally not expressed.

Discussion

In this study, we report that the neurite growth-promoting and degeneration-attenuating effects of a novel *Lymnaea* protein, named caltubin, can be transferred to rodent neurons. Expression of caltubin in rat PC12 cells or mouse cortical neurons effectively enhanced outgrowth and attenuated axonal degeneration following transection. These findings implicate caltubin as a novel molecular player in a conserved cellular mechanism of neurite outgrowth and highlight its potential as a possible therapeutic tool in nerve regeneration.

Local protein synthesis is one of the crucial mechanisms regulating neurite outgrowth, growth cone behavior, and axonal regeneration (Gumy et al., 2010). Whereas the exact downstream mechanisms by which local protein synthesis regulates axon growth remain unclear, most mRNAs that are identified in axonal growth cones appear to be structural proteins in both invertebrates (Plunet et al., 2002; Gioio et al., 2004) and in vertebrates (Willis et al., 2005; Hengst et al., 2009), and translational regulation of these proteins occurs during axon regeneration (Twiss et al., 2000). For instance, stimulus-driven axonal growth is dependent on local translation of the cytoskeleton regulator PAR (Hengst et al., 2009), and inhibition of local translation of the actin regulating protein β -thymosin enhances rates of neurite outgrowth (van Kesteren et al., 2006). In this study, we provided multiple lines of evidence that local protein synthesis of caltubin is required for neurite regeneration through regulation of β -tubulin level. First, *caltubin* mRNA and protein are highly expressed in growth cones of snail neurons. Second, local caltubin levels in isolated neurites can be reduced by targeted siRNA knockdown or anisomycin treatment. Third, reduction of local caltubin levels in isolated neurites suppresses neurite extension and regeneration following injury in cultured CNS neurons. Fourth, caltubin protein level increased in distal neurites 3 h after transection, indicating the potential *de novo* synthesis of proteins in these regions. Finally, caltubin colocalizes and physically interacts with tubulin, and a reduction of local levels of caltubin by targeted siRNA knockdown leads to a decrease in local tubulin levels. Together, our findings indicate that local synthesis of *de novo* proteins is essential for neurite regeneration during this period of time, and that locally synthesized caltubin promotes regenerative neurite outgrowth.

The cytoskeleton is an essential element for cell architecture and neurite structure. Most of the local transcripts are cytoskeletal proteins (e.g., microtubule, microfilament, and intermediate filament proteins) (Kaplan et al., 1992; Eng et al., 1999; Vogelaar et al., 2009), or proteins that regulate cytoskeletal dynamics (e.g., Rho and β -thymosin) (Campbell et al., 2001; Verma et al., 2005; van Kesteren et al., 2006). Stabilization and continuous restructuring of the cytoskeleton plays an important role in the processes of neurite formation and regeneration, allowing growth cone formation, vesicle accumulation, and neurite elongation (Spira et al., 2001; Witte et al., 2008). We found that caltubin colocalizes with and binds to both α - and β -tubulin. Thus, the neurite growth-promoting effects of caltubin are likely through local regulation of tubulin. Caltubin knockdown decreased local α - and β -tubulin protein levels, but did not affect β -actin or NCS-1 levels, indicating that caltubin specifically regulates tubulin levels to achieve its effects on promoting neurite outgrowth and regeneration. Microtubules are comprised of $\alpha\beta$ -tubulin heterodimers (Westermann and Weber, 2003), and polymerization of microtubules stabilizes the cytoskeleton (Nakano et al., 2010), leading to maintenance of neurite integrity (Karima et al., 2010), and promotion of neurite outgrowth (Tanaka et al., 1995) and regeneration (Ertürk et al., 2007). While the mechanisms by which caltubin may regulate tubulin levels remain unclear, we speculate that binding of caltubin may cause overall stabilization of tubulin polymers, which in turn enhances neurite elongation. Similarly, the ring finger protein ZNRF1, encoded by a nerve injury-induced neuronal gene *Nin283* (Araki et al., 2001), interacts with β -tubulin to enhance neurite-like elongation (Yoshida et al., 2009), suggesting that other tubulin-binding proteins may also regulate axonal elongation following injury. Knockdown of caltubin may destabilize microtubule proteins, leading to neurite

retraction. Alternatively, caltubin may regulate local translation or degradation of tubulin in the axon. While further investigation is required for understanding how exactly caltubin regulates tubulin levels, our data clearly indicate that in *Lymnaea* neurons caltubin binding to tubulin is a prerequisite for efficient axon growth and regeneration.

Vertebrate and invertebrate growth cones and axons share many common mechanisms of neuronal growth, including organization of the microtubule cytoskeleton, response to axon guidance molecules, and dynamic changes in intracellular calcium signaling (Dent et al., 2003). Many of the crucial proteins involved in neurite outgrowth and regeneration, including tubulin, are highly conserved across species (Burns and Surridge, 1990). Yet vertebrate and invertebrate adult central neurons differ in their regenerative capacity after injury: in most vertebrates, central neurons seldom regenerate injured neurites, whereas injury to invertebrate central neurons results in a strong regenerative response. Taking advantage of the high conservation between tubulins across species, we tested whether caltubin, which has no obvious sequence orthologs in mammals, might also promote neurite outgrowth and regeneration in mammalian cells. We show that caltubin not only binds to endogenous tubulin but also promotes NGF-induced axonal outgrowth in PC12 cells and cultured mouse cortical neurons, and reduces rates of axonal retraction. Thus, our data suggest that caltubin might be one of the critical molecules essential for the robust intrinsic ability to regenerate in snail neurons, and that some of its downstream mechanisms are conserved in mammalian central neurons. Indeed, our findings show that the molecular interactions and cellular functions of caltubin are conserved in invertebrate and vertebrate neurons, leading to a promising new avenue for using an invertebrate protein as a potential therapeutic target to remodel neuronal properties favoring axonal regeneration and/or recovery from neural injury and neurodegenerative disorders.

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