Neurite outgrowth is mediated by the exocytosis of intracellular vesicles at the tips of elongating neuronal processes. The lysosomal vesicle-associated soluble N-ethylmaleimide-sensitive factor attachment protein receptor tetanus neurotoxin insensitive vesicle-associated membrane protein (TI-VAMP)/VAMP7 was previously implicated in membrane fusion events mediating neurite outgrowth, but the participation of lysosomes in this exocytic process has remained unclear. Here, we show that VAMP7 and the lysosomal glycoprotein Lamp1 extensively colocalize in vesicles present throughout the soma and neurite outgrowths of primary sympathetic neurons. Synaptotagmin VII (Syt VII), a Ca\(^{2+}\)-sensing synaptotagmin isoform previously shown to interact with VAMP7 during lysosomal exocytosis in fibroblasts, was detected on a subset of these lysosomal glycoprotein 1 (Lamp1)/VAMP7-positive neuronal vesicles. Ionophore-stimulated exocytosis triggered exposure of the luminal domains of both Lamp1 and Syt VII at overlapping sites on the neuronal surface, indicating that the Syt VII-containing lysosomal compartments fuse with the plasma membrane in response to [Ca\(^{2+}\)]\(_i\) elevation. To determine whether Syt VII was required for the exocytic events mediating neurite extension, we followed the development of superior cervical ganglion neurons explanted from Syt VII-deficient mice. The results revealed a marked defect in neurite outgrowth and arborization, suggesting that Ca\(^{2+}\)-dependent, Syt VII-regulated exocytosis of late endosomes/lysosomes plays a role in the addition of new membrane to developing neurite extensions.

Key words: calcium; dendrite; development; exocytosis; fluorescence microscopy; knock-out mice

Introduction

The extension, branching, and interconnection of axons and dendrites play a central role in development of the nervous system (Prochiantz, 1995). Regrowth of neuritic trees is also observed during the regeneration of injured neurons, a process that is modulated by the influx of Ca\(^{2+}\) at the injury site (Rehder et al., 1992; Chu and Tator, 2001). Consistent with a mechanism based on regulated exocytosis, neurite outgrowth involves the polarized addition of new membrane at the tips of elongating processes (Craig et al., 1995; Futerman and Banker, 1996; Bradke and Dotti, 1997; Zakharenko and Popov, 1998; Martinez-Arca et al., 2001).

Vesicle-associated soluble N-ethylmaleimide-sensitive factor attachment protein receptor (SNARE) molecules present on intracellular vesicles (v-SNAREs) and target membranes (t-SNAREs) are essential components of the molecular machinery responsible for membrane fusion (Rothman, 1994). Recent studies implicated the tetanus neurotoxin insensitive v-SNARE (TI-VAMP)/VAMP7 in the exocytic events responsible for neurite extension. (Martinez-Arca et al., 2000). The NH\(_2\)-terminal longin domain of VAMP7 inhibited SNARE complex formation and exerted dominant-negative effects on both axonal and dendritic outgrowth when introduced into neuroendocrine PC12 cells or primary neurons in culture (Martinez-Arca et al., 2000, 2001). Inhibition in neurite outgrowth was also observed after RNA interference-mediated silencing of VAMP7 (Alberts et al., 2003). These observations implicated VAMP7-containing intracellular compartments as a source of membrane for neurite outgrowth. However, the intracellular signaling pathways regulating VAMP7-dependent vesicular exocytosis and the nature of the VAMP7-containing compartments in neurons remain poorly understood.

VAMP7 was originally identified as a synaptobrevin-related SNARE (D’Esposito et al., 1996) resistant to cleavage by Tetanus and Botulinum neurotoxins (Galli et al., 1998). In fibroblasts, VAMP7 is localized on vesicular compartments containing the late endosomal/lysosomal glycoprotein 1 (Lamp1) (Advani et al., 1999; Rao et al., 2004) and mediates membrane traffic between late endosomes and lysosomes (Advani et al., 1999). Recently, VAMP7 was also shown to be required for Ca\(^{2+}\)-triggered lyso-
somal exocytosis in fibroblasts (Rao et al., 2004). After elevation in intracellular free Ca$^{2+}$ ([Ca$^{2+}]_i$), VAMP7 forms SDS-resistant complexes with syntaxin 4 and soluble N-ethyl maleimide-sensitive factor attachment protein 23 (SNAP23), two plasma membrane t-SNAREs that interact in a Ca$^{2+}$-dependent manner with the lysosomal synaptotagmin VII (Syt VII) (Rao et al., 2004). Syt VII is an evolutionarily conserved member of the synaptotagmin family of Ca$^{2+}$ sensors (Andrews and Chakrabarti, 2005). Two Ca$^{2+}$-binding C2 domains present on the cytosolic region of synaptotagmins mediate Ca$^{2+}$-dependent interactions with membrane phospholipids and SNARE complex components, facilitating membrane fusion (Tucker and Chapman, 2002). Previous studies from our laboratory showed that Syt VII regulates Ca$^{2+}$-triggered lysosomal exocytosis (Martinez et al., 2000) and the rescaling of plasma membrane wounds (Reddy et al., 2001). The functional association of Syt VII with VAMP7 observed during lysosomal exocytosis in fibroblasts (Rao et al., 2004) suggested that similar Ca$^{2+}$-regulated exocytic events might be involved in neurite outgrowth. We investigated this possibility using primary neurons isolated from mouse sympathetic cervical ganglia (SCGs). Our results show that VAMP7 and Syt VII colocalize in peripheral late endosomal/lysosomal compartments that exocytose after [Ca$^{2+}]_i$ elevation, and that genetic ablation of Syt VII results in defective neurite outgrowth.

Materials and Methods

Materials. Monoclonal anti-mouse Lamp1 antibodies (1D4B) were obtained from the Developmental Studies Hybridoma Bank (Iowa City, IA), and monoclonal anti-VAMP2 (C6E9.1) was obtained from Synaptic Systems (Goettingen, Germany). Polyclonal rabbit antibodies against the luminal NH2-terminal sequence of synaptotagmin VII were generated and affinity-purified as described previously (Martinez et al., 2000). Monoclonal antibodies (mAbs) (clone 158.2) against Ti-VAMP/VAMP7 were a generous gift from Dr. Thierry Galli (Institut National de la Sante et de la Recherche Medicale U536, Institut du Fer-a-Moulin, Paris, France). Fluorescent secondary Alexa Fluor 488 or 546 antibodies were purchased from Invitrogen (Carlsbad, CA), and ionomycin was purchased from Calbiochem (San Diego, CA). Mouse nerve growth factor (NGF 2, 5S) was purchased from Alomone Labs (Jerusalem, Israel), and arabinoside C was purchased from Sigma (St. Louis, MO).

Immunofluorescence. Localization of antigens of interest were performed on monolayers fixed with 2% paraformaldehyde (PFA). After permeabilization with 0.1% saponin for 15 min, the cells were incubated with 1:100 dilutions of antibodies against LAMP1, Ti-VAMP/VAMP7, and VAMP2, followed by the appropriate fluorescent secondary antibodies. For staining with antibodies against synaptotagmin VII (1:100 dilution), cells were permeabilized with 0.1% Triton X-100. Double immunohistochemistry for observation of colocalization of the markers was performed with an initial 0.1% saponin permeabilization in the case Lamp1 staining, followed by permeabilization in 0.05% Triton X-100 for Ti-VAMP/VAMP7 or Syt VII staining. Secondary antibodies alone were used as specificity controls and uniformly resulted in very low background levels of fluorescence.

Detection of surface Lamp1 and Syt VII after exocytosis. Neuronal cultures were washed and incubated at 37°C with 0.5 ml of either PBS alone or PBS containing 5 μM ionomycin for 3–5 min. The cells were then incubated at 4°C for 30 min with anti-LAMP1 (1D4B hybridoma culture supernatant) or anti-Syt VII luminal domain antibodies. Cells were then fixed with 2% PFA for 15 min at 4°C, washed in PBS, and incubated with the appropriate secondary antibodies for 30 min at room temperature. Nuclei were stained with 4′,6′-diamidino-2-phenylindole (DAPI) (Sigma).

SCG explant assay. Syt VII+/+ and Syt VII−/− mice were generated as described previously (Chakrabarti et al., 2003) and back-crossed for eight generations into the C57BL/6 background. Superior SCGs were bilaterally dissected from newborn (1–2 d of age) Syt VII+/+ or Syt VII−/− mice, washed in ice-cold PBS, and explanted to Matrigel-coated 35 mm glass-bottom culture dishes (Matake Corporation, Ashland, MA). A drop of cold Matrigel (BD Biosciences, PharMingen, San Diego CA) diluted 1:4 in DMEM was placed at the bottom of the dish and incubated at room temperature until the gel solidified. Each dissected SCG cleaned of all surrounding connective tissue was deposited on the top of the drop, additional 20 μl of diluted Matrigel was added, and the dish was incubated for 15 min at room temperature. DMEM (1.5 ml) 10% FCS supplemented with 50 ng/ml NGF (Alomone Labs) was then added, and the explants were incubated for 3, 12, 24, 72, and 120 h at 37°C 5% CO2. At each time point, phase contrast images of SCG explants were acquired using a 10× objective (100× final magnification) of a Zeiss (Thornwood, NY) Axiovert 135 microscope equipped with a Hamamatsu (Shizuoka, Japan) Orca II cooled CCD camera controlled by MetaMorph software (Molecular Devices, Sunnyvale, CA).

Explant outgrowth measurements. For the early time points (3–24 h), explants were small enough to be visualized and measured in a single microscopic field. After this initial growth period, four separate images covering the complete ganglion were taken from each explant, and the outgrowth length was determined on scanned images of each quadrant captured with a 10× objective. Measurements were made in exported eight-bit TIFF files using the image analysis software MetaMorph. Maximum neurite length was determined as follows: the MetaMorph “draw” function was used to draw a straight line from the border of the explants to the tip of the longest neurite or neurite bundle at 30 different locations covering each of the four quadrants of explants, as described previously (Arantes et al., 2003). Maximum neurite length was defined as the average of these 30 measurements, and the values were calibrated in micrometers using the MetaMorph “calibrate” function. In each experiment, the minimum number of ganglia measured per time point was three, and experiments were repeated three to five times. The results were expressed as the average and SD of the results of each separate experiment and analyzed for statistical significance with an unpaired Student’s t test (p values <0.05 were considered significant).

Isolated neuron cultures. Primary neuronal cultures were prepared from SCGs of newborn Syt VII+/+ or Syt VII−/− mice, as described previously (Arantes et al., 2000). SCGs were bilaterally dissected from newborn (1–2 d of age) C57BL/6 mice (average of 16 ganglia per litter), washed in PBS, and combined in a single pool. The cells were incubated for 1 h at 37°C in 0.25% trypsin solution (Invitrogen, Carlsbad, CA) followed by mechanical dissociation with a long-tip, fire-polished Pasteur pipette until individual cells could be visualized and counted in a Neubauer chamber. Cells were plated at 104/dish on glass-bottom 35 mm dishes previously coated with Matrigel diluted 1:4 in DMEM (Invitrogen) and grown in DMEM 10% FCS containing 50 ng/ml NGF (re supplied by the Developmental Studies Hybridoma Bank). After the growth of glia and other accessory cells) for the entire experimental period. Cultures were kept at 37°C in 5% CO2 for 1, 2, 5, 4, and 24 h, followed by fixation in 2% PFA for 30 min at room temperature. Quantification of neurite outgrowth in isolated neurons. Images of randomly chosen fields of fixed SCG isolated neuron cultures were obtained under 1000× magnification using a Zeiss Axiosvert 135 microscope equipped with a Hamamatsu Orca II cooled CCD camera and MetaMorph software. Images from at least 60 individual fields were captured, resulting in the analysis of 50–150 neurons per time point, per experiment. Neurons were distinguished from other cell types by immunofluorescence with antibodies against VAMP2. Quantification of neurite length was done on exported eight-bit TIFF files using the MetaMorph image analysis software. Neurites were outlined using the draw function, and their perimeter was automatically measured and calibrated in micrometers using the MetaMorph calibrate function. The number of primary neurites (defined as the segments of neurites originating directly from the cell body of neurons) and the number of branching points were microscopically counted. For each microscopic field, the number of neuronal cell bodies was also counted, and the data were expressed as the average neurite length/neuron, number of primary neurites/neuron, and branching points/neuron. Because at 24 h the neurite network was too complex to allow accurate branch point measurements, at this time point, neurite density was determined by superposing two vertical lines.
on the captured images, as described previously (Arantes et al., 2000; Ronn et al., 2000). Neurite density was defined as the average number of neurites that were intersected by the lines. The results were expressed as the average and SD of three independent experiments and analyzed for statistical significance with an unpaired Student’s t test (p values <0.05 were considered significant).

Results
Late endosomes/lysosomes containing Lamp1 and VAMP7 are present in neurite outgrowths, including the tips of elongating processes

Previous studies concluded that the late endosomes and lysosomes of primary neurons were predominantly located in the soma and proximal dendrites (Parton et al., 1992; Craig and Banker, 1994). Subsequent work using more sensitive immunofluorescence techniques revealed the presence of lysosomes throughout axon shafts, including branching points (Overly and Hollenbeck, 1996). However, it remained unclear whether late endosomes/lysosomes were present at sites of new membrane addition, such as the tips of elongating neurites during neuronal development. To investigate this issue, we performed immunofluorescence with antibodies against the lysosomal marker Lamp1 on SCG neurons at increasing time periods after plating dissociated ganglia in the presence of NGF. After 1–2 h, the culture contained multiple clusters of neurons (identified by labeling with anti-synaptotagmin I and anti-VAMP2 antibodies) (Fig. 2B), some of which contained short neurite extensions (Fig. 1A). Although Lamp1-positive compartments were abundant in cell bodies, clusters of vesicles reactive with the antibody were also detected at distal regions of the elongating neurites (Fig. 1B). The presence of lysosomes within growing neurites was more evident after 4 h of culture, when Lamp1-positive vesicles were frequently observed at branching points and at the tips of elongating processes (Fig. 1C–E).

Previous studies demonstrated a role for the late endosomal v-SNARE VAMP7 in neurite outgrowth from PC12 cells and primary hippocampal neurons (Martinez-Arca et al., 2000, 2001; Alberts et al., 2003). Although VAMP7 was found to colocalize with the late endosomal/lysosomal marker CD63 (Coco et al., 1999), the late endosomal/lysosomal nature of the VAMP7 compartments was questioned in those studies because of an apparent lack of VAMP7 colocalization with Lamp1 (Coco et al., 1999). To clarify this issue, we performed double immunofluorescence assays with antibodies against Lamp1 and VAMP7 on developing SCG neurons. We detected extensive colocalization between these two markers, both at early (Fig. 2A, 1 h) and later (Fig. 2A, 4 h) time points. Vesicles immunoreactive for both Lamp1 and VAMP7 were present throughout the neurons, and clusters of vesicles labeled by both antibodies were frequently observed within growth cones (Fig. 2A, 4 h). The synaptic vesicle SNARE VAMP2 was also detected within growth cone regions, but its localization was clearly distinct from the lysosomal marker Lamp1 (Fig. 2B). Thus, these results demonstrate that in primary sympathetic neurons, VAMP7-containing compartments completely overlap with Lamp1-positive late endosomes/lysosomes, as shown previously in other cell types (Advani et al., 1999; Rao et al., 2004).

The lysosomal synaptotagmin Syt VII is present in Lamp1-positive compartments of SCG neurons and translocated to the cell surface after Ca2+-triggered exocytosis
Syt VII was previously shown to colocalize with lysosomal markers in fibroblasts (Martinez et al., 2000; Caler et al., 2001; Jaiswal et al., 2002; Chakrabarti et al., 2003), epithelial cells (Caler et al., 2001), keratinocytes (Hakansson et al., 2005), and the neuroendocrine cell line PC12 (Fukuda et al., 2004; Wang et al., 2005). To determine whether the same occurred in primary neurons, we performed immunolocalization of Syt VII in isolated SCG neurons. These assays revealed the presence of intracellular vesicular compartments containing both Lamp1 and Syt VII, within the cell bodies and also at peripheral regions associated with developing neurite extensions (Fig. 3). The relative abundance of Lamp1 and Syt VII on the double-positive vesicles detected varied, suggesting the existence of Syt VII or Lamp1-enriched domains on the same compartments.

Figure 1. Lamp1-containing vesicles are present in the cell body and throughout neuritic outgrowths of SCG neurons. A, DIC images of SCG neurons at 2.5 h (A) or 4 h (C) after plating, showing the typical pyramidal, globular shape of the cells with thin extensions originating from the soma (arrows). B, D, E, Immunofluorescence of saponin-permeabilized SCG neurons with anti-Lamp1 mAbs (green) and nuclear DAPI staining (blue) after 1 h (B) and 4 h (D, E) in culture. Arrows point to distal regions and branching points of growing neurites, where Lamp1-positive vesicles were detected (D, E).
Previous studies from our laboratory showed that the luminal domains of Lamp1 and Syt VII are exposed on the surface of fibroblasts, when lysosomal exocytosis is induced by elevation in \([\text{Ca}^{2+}/\text{H}^{11001}]\) (Reddy et al., 2001). To determine whether the lysosomal exocytic pathway was functional in SCG neurons, we exposed cultures of developing neurons to the \(\text{Ca}^{2+}/\text{H}^{11001}\) ionophore ionomycin and performed immunofluorescent detection of surface-exposed Lamp1 and Syt VII. As reported previously for fibroblasts (Rodriguez et al., 1997; Reddy et al., 2001), elevation in \([\text{Ca}^{2+}/\text{H}^{11001}]\) resulted in a punctate labeling pattern on the surface of SCG neurons, when intact cells were surface-stained with antibodies against the luminal domain of Lamp1 (Fig. 4). Several of the surface-exposed, Lamp1-positive puncta colocalized precisely with areas where the luminal domain of Syt VII was also detected with specific antibodies. Spots containing both Lamp1 and Syt VII were detected predominantly at the periphery of developing neurons and, in some instances, appeared to be associated with sites of neurite outgrowth (Fig. 4). Consistent with the fact that not all Lamp1-positive intracellular compartments in SCG neurons contain detectable Syt VII (Fig. 3), a fraction of the surface-exposed luminal domain of Lamp1 was not associated with Syt VII and was detected in a more perinuclear location (Fig. 4). These observations suggested that lysosomal compartments of sympathetic neurons are capable of fusing with the plasma membrane after \([\text{Ca}^{2+}/\text{H}^{11001}]\) elevation, and that a more peripheral fraction of these Lamp1-positive exocytic compartments contains the \(\text{Ca}^{2+}\) sensor Syt VII.

**Neurite outgrowth is impaired in SCG neurons from Syt VII-deficient mice**

Previous studies in fibroblasts demonstrated that genetic ablation of Syt VII inhibits lysosomal exocytosis, when the process is triggered by \(\text{Ca}^{2+}/\text{H}^{11001}\) influx at sites of membrane injury (Chakrabarti et al., 2003; Roy et al., 2004) or by mobilization of intracellular \(\text{Ca}^{2+}\) stores through receptor-mediated agonists (Caler et al., 2001; Jaiswal et al., 2004). In contrast, the higher and more widespread \([\text{Ca}^{2+}/\text{H}^{11001}]\) elevation resulting from stimulation with \(\text{Ca}^{2+}/\text{H}^{11001}\) ionophores can also trigger lysosomal exocytosis in Syt VII-deficient cells, albeit with a markedly altered secretory pattern when compared with wild-type cells (Jaiswal et al., 2004). However, concerns about toxic and ATP-depleting effects of \(\text{Ca}^{2+}/\text{H}^{11001}\) ionophores has limited the usefulness of this procedure for the study of physiologically relevant exocytosis (Andrews and Chakrabarti, 2005). For this reason, we examined the consequences of Syt VII ablation using physiological assays of NGF-dependent neurite outgrowth, a process already shown to require the exocytosis of intracellular compartments containing VAMP7 (Martinez-Arca et al., 2000, 2001).

SCG ganglia isolated from Syt VII\(/+\) or Syt VII\(/-/-\) mice were cultured within Matrigel matrices in the presence of NGF for increasing periods of time, and the average length of neurites outgrowing from the explants was determined on microscopically acquired images. In independent experiments, a reduction in neurite outgrowth was evident in ganglion explants from Syt VII\(/-/-\) mice after 12–120 h of culture (Fig. 5A, B).

To characterize more precisely the apparent defect in neurite outgrowth resulting from Syt VII deficiency, we examined isolated SCG neurons during their initial 24 h of development in culture. Isolated ganglia were dissociated, and the isolated neurons were plated in Matrigel-coated dishes in the presence of...
NGF and fixed and imaged after 1, 2.5, 4, and 24 h. Although there was a reduction in the number of neurons remaining attached to the substrate between 4 and 24 h, the loss was similar in Syt VII/H11001/H11001 and Syt VII/H11002/H11002 cultures (supplemental Fig. 1, available at www.jneurosci.org as supplemental material). The images were analyzed for average neurite length (total area occupied by extensions emanating from the cell bodies), number of primary neurites, and number of branching points (Figs. 6, 7). Statistically significant differences were observed between Syt VII/H11001/H11001 and Syt VII/H11002/H11002 neurons in three independent experiments. The average neurite length was markedly reduced in Syt VII/H11002/H11002 neurons after 24 h in culture (Fig. 7A), and significant decreases in the number of primary neurites and in the number of branching points were also detected after 1, 2.5, 4, and 24 h (Fig. 7B–D).

Together, these results indicate that the late endosomal/lysosomal Ca\(^{2+}\) sensor Syt VII participates in the exocytic process mediating neurite outgrowth. Syt VII absence from SCG neurons results in a defect in new membrane deposition and in a markedly reduced complexity of the arborification process of developing growth cones.

Discussion

Previous studies demonstrated that intracellular compartments containing the v-SNARE VAMP7 participate in the exocytic delivery of intracellular membrane to neurite extension sites (Martinez-Arca et al., 2000, 2001; Alberts et al., 2003). It remained unclear, however, which were the properties of these VAMP7-positive compartments that make them responsive to signals regulating localized exocytosis. In this study, we show that VAMP7 and the lysosomal marker Lamp1 fully overlap in primary sympathetic SCG neurons. These VAMP7/Lamp1-positive late endosomes/lysosomes are also present within growth cones at distal regions of developing neuronal processes. A subset of these vesicles contains Syt VII, the ubiquitously expressed synaptotagmin isoform previously shown to regulate Ca\(^{2+}\)-triggered exocytosis. We found that Syt VII/VAMP7-positive vesicles in SCG neurons behave as Ca\(^{2+}\)-regulated secretory vesicles, fusing with the plasma membrane after \([\text{Ca}^{2+}]_\text{i}\) elevation. These findings suggested that Syt VII-dependent Ca\(^{2+}\)-triggered exocytosis might be involved in the mobilization of late endosomes/lysosomes to the plasma membrane during neurite outgrowth. We examined this question by following the development in culture of SCG neurons isolated from wild-type or Syt VII-deficient mice. Our results show that the number, length,
and complexity of neuritic tree outgrowths are significantly reduced in Syt VII-deficient neurons, directly implicating the Ca\(^{2+}\) sensor Syt VII in this process.

Several studies have suggested a role for [Ca\(^{2+}\)]\(_i\) elevation as a trigger for neurite outgrowth. In addition to being required for membrane resealing in injured neurites, Ca\(^{2+}\) influx induces the formation of growth cones (Davenport and Kater, 1992; Rehder et al., 1992; Ziv and Spira, 1997; Chu and Tator, 2001; Shoemaker et al., 2005). A few minutes after membrane injury or the local application of Ca\(^{2+}\) ionophores, *Aplysia* giant neuron axonal segments dedifferentiate into growth cones and generate a vigorous neuritic tree outgrowth (Ziv and Spira, 1997). Similar observations were made in rat SCG neurons; transected neurites were able to regrow in the presence of extracellular Ca\(^{2+}\) but not when neurons were preloaded with the Ca\(^{2+}\) chelator BAPTA-AM (Chu and Tator, 2001). In our experimental system, mouse SCG neurons were cultured in the presence of NGF, which is essential for their long-term survival. NGF exerts its neurotrophic action through Trk receptors (Kalb, 2005), which when activated can lead to small and rapid increases in [Ca\(^{2+}\)]\(_i\) (Pandiella-Alonso et al., 1986; Jiang and Guroff, 1997). Basal [Ca\(^{2+}\)]\(_i\) levels seem to be important for NGF-dependent survival of sympathetic neurons in culture (Koike and Tanaka, 1991), but it remains unclear whether NGF-dependent Ca\(^{2+}\) signaling is required for neurite outgrowth (Tolkovsky et al., 1990). Although the neuritic tree outgrowth observed in our studies may have been, at least in part, triggered by Ca\(^{2+}\) influx through wounds on neurite membranes generated during SCG isolation, a role for NGF-dependent [Ca\(^{2+}\)]\(_i\) increases cannot be ruled out. Because NGF is required for long-term survival of SCG neurons in culture, it has not been possible to directly examine the role of NGF as a mediator of Ca\(^{2+}\) signaling and of Syt VII-dependent neurite extension.

Axonal growth and dendrite extension are critical events in the development of the nervous system. Despite the marked defect observed in neurite outgrowth from explanted SCG neurons in culture, Syt VII deficiency does not result in abnormal brain development or in any obvious neurological phenotype (Chakrabarti et al., 2003). A similar lack of a brain development phenotype was observed in SNAP25 mouse null mutants (Washbourne et al., 2002), in contrast to the strong inhibition in neurite outgrowth seen in isolated hippocampal neurons after SNAP25 cleavage with Botulinum neurotoxin A (Grosse et al., 1999). The lack of a direct correlation between *in vitro* and *in vivo* phenotypes is often related to the existence of functionally redundant pathways. This could have occurred at the level of the molecules mediating vesicular traffic (such as additional synaptotagmin isoforms in Syt VII/−/− mice, or the functional replacement of SNAP25 by SNAP23) or at the level of the signaling pathways triggering neurite extension. One example of a redundant signaling pathway is the inhibition in neurite outgrowth observed in Ryk-deficient dorsal root ganglion (DRG) neurons stimulated with Wnt, but not with NGF, providing an explanation for the apparently normal *in vivo* DRG axon outgrowth of Ryk-deficient mice (Lu et al., 2004). It will be of interest to determine whether...

![Figure 5](image-url) Neurite outgrowth is impaired in SCG explants from Syt VII-deficient mice. A, Average neurite length after increasing periods of time in culture. The data are expressed as the mean ± SD of neurite length average values from independent experiments (n = 3–5). B, Phase contrast images of SCG explants after 24 and 120 h in culture, illustrating the reduced neurite outgrowth observed in Syt VII/−/− neurons (right panels).

![Figure 6](image-url) Neurite growth is impaired in neurons isolated from SCG explants from Syt VII-deficient mice. Phase contrast images of isolated SCG neurons after increasing periods of time in culture show the reduced complexity of the outgrowing neuritic trees in Syt VII/−/− neurons.
the impaired neuritic outgrowth we observed in cultured Syt VII-deficient SCG neurons has an impact on the postnatal maintenance of dendritic arbor structure in vivo. Recent evidence suggests that dendrite branch maintenance may be essential for sustaining normal brain function (Zoghbi, 2003; Moresco et al., 2005).

References


Davenport RW, Kater SB (1992) Local increases in intracellular calcium have an impact on the postnatal maintenance of dendritic arbor structure in vivo. Recent evidence suggests that dendrite branch maintenance may be essential for sustaining normal brain function (Zoghbi, 2003; Moresco et al., 2005).

Figure 7. Neurite extension and arborization is impaired in neurons isolated from SCG explants from Syt VII-deficient mice. A, Average neurite length after increasing periods of time in culture. B, Average number of primary neurites after increasing periods of time in culture. C, Average number of branching points after increasing periods of time in culture. D, Average neurite density after 24 h in culture. The data are expressed as the mean ± SD of average values from independent experiments (n = 3).


