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Journal Article

Kyoto University
An evolutionarily conserved protein CHORD regulates scaling of dendritic arbors with body size

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Most organs scale proportionally with body size through regulation of individual cell size and/or cell number. Here we addressed how postmitotic and morphologically complex cells such as neurons scale with the body size by using the dendritic arbor of one Drosophila sensory neuron as an assay system. In small adults eclosed under a limited-nutrition condition, the wild-type neuron preserved the branching complexity of the arbor, but scaled down the entire arbor, making a “miniature”. In contrast, mutant neurons for the Insulin/IGF signaling (IIS) or TORC1 pathway exhibited “undergrowth”, which was characterized by decreases in both the branching complexity and the arbor size, despite a normal diet. These contrasting phenotypes hinted that a novel regulatory mechanism contributes to the dendritic scaling in wild-type neurons. Indeed, we isolated a mutation in the gene CHORD/morgana that uncoupled the neuron size and the body size: CHORD mutant neurons generated miniature dendritic arbors regardless of the body size. CHORD encodes an evolutionarily conserved co-chaperone of HSP90. Our results support the notion that dendritic growth and branching are controlled by partly separate mechanisms. The IIS/TORC1 pathways control both growth and branching to avert underdevelopment, whereas CHORD together with TORC2 realizes proportional scaling of the entire arbor.

The organ size is controlled through changes in rates of cell growth and division

The underlying mechanisms of organ size control, such as Insulin/IGF signaling (IIS) and Target of Rapamycin complex 1 (TORC1) signaling pathways, have been intensively investigated by using proliferating and morphologically simple cells, such as those in epithelial monolayers of the Drosophila imaginal disc. However, it is still largely unknown how the size of postmitotic and morphologically complex cells such as neurons scale with body size and which molecules control such scaling. We therefore explored these scaling mechanisms in Drosophila neurons.

For our investigation, we used the dendritic arbor of one class of sensory neurons in Drosophila adults as a readout (Figure 1A and 1B). Dendrites are the antennae of neurons that receive and integrate sensory and/or synaptic inputs. Our model neuron in this study, previously designated as v'ada, is one of the dendritic arborization (da) neurons, whose arbor for adult life is regenerated during pupal stages and entirely covers the lateral plate (pleura) in the abdomen (Figure 1A–1C)

Results

Scaling of dendritic arbors of the wild-type da neuron with body size. To examine how body-size changes affect the size and the branching pattern of the dendritic arbor of the wild-type neuron, we starved larvae beyond 91–96 hr after egg laying (AEL). This is because it has been shown that larvae exposed to such starvation after the “critical weight” stop body growth, but develop to become fertile adults that are smaller than normal, without any developmental delay. Under this mild starvation condition, the size of the dendritic arbor was significantly decreased in proportion to the decreased body size (Figure 1D–1G). Importantly, the number of branch endings was not changed and the branch density (total length/arbor size and ending number/arbor size) was significantly increased (Figure 1H–1K). We also quantified the branching complexity, by assigning Strahler orders to individual segments (intervals between branching points; see Figure 1L). The numbers of individual order segments were similar under the fed and starved conditions (Figure 1M), while 2nd to 4th order segments became shorter under the starved condition (Figure 1N). These results are consistent with the hypothesis that the
wild-type neuron can respond commensurately to the decrease in body size and/or the starvation and can form a “miniature” dendritic arbor by tuning its dendritic segment length.

Neurons with defective IIS/TORC1 signaling pathways show dendritic “undergrowth.” The above result suggested that the wild-type neuron is able to scale with the body size and/or the nutrition condition, while keeping the branching complexity intact. We wondered if the IIS/TORC1 pathways (Supplementary Figure S1A) participate in this scaling, and whether defects in the pathways would affect the size and the branching pattern of the dendritic arbor in adults, when larvae were raised under the normal food condition.

Neurons with disruptions of Drosophila insulin receptor (dinr) or Akt downsized and simplified the dendritic arbors, as evidenced by decreases in the arbor size, the ending number, and the total length (Supplementary Figure S1B–S1D and S1H–S1J). In contrast, the branch density (total length/arbor size) and ending number/arbor size (K) was not significantly altered from that of the wild-type neuron (Supplementary Figure S1K and S1L). A loss of function mutation of tor, overexpression of a dominant negative form of Phosphoinositide 3-kinase (PI3K), or a knockdown of raptor encoding an essential component of TORC1 not only decreased the arbor size, but also the branching complexity (Supplementary Figure S1E–S1G). Thus this “undergrowth” phenotype is distinct
from the “miniature” dendrite of the wild-type neuron under the starved condition (Figure 2L), suggesting that a regulatory mechanism other than the IIS/TORC1 pathways may contribute to the dendritic scaling in the normal da neuron.

CHORD mutant neurons form miniature dendrites. To explore the hypothetical mechanism of the dendritic scaling, we conducted a forward genetic screen by employing the mosaic analysis with a repressible cell marker (MARCM) system. To facilitate the generation of mosaic clones, we made “SOP-FLP” lines that express FLP recombinase in sensory organ precursors (SOPs; see details in Methods). In our screening under the fed condition, we found a mutant chromosome that generated miniature dendritic patterns in homozygous neurons (Figure 2B and 2C). Indeed a number of quantitative features indicated that the arbor of this mutant neuron was a proportionally scaled down miniature or a microcopy of the wild-type neuron (Figure 2E–2I): decreases in the arbor size and the total dendritic length, an unaltered ending number, an increase in branch density, profiles of the Strahler-order analysis similar to those of the wild-type neuron under the starvation condition (compare Figure 1M–1N with Figure 2J–2K), and an unaltered distribution of angles at individual branching points (data not shown). In a word, the mutant neuron reduced the dendritic segment length (Figure 2K), without simplifying the branching pattern, which is reminiscent of how the wild-type neuron constructed its arbor under the starved condition (Figure 1F).

Figure 2 | CHORD mutant neurons form “miniature” dendrites. (A) Whole-genome sequencing identified a 1 bp deletion in the CHORD gene unique to the mutant strain that was isolated from our forward genetic screening (bottom). We denoted this allele as CHORD2 hereafter. Note that we determined the sequences of flies heterozygous for the mutation. (Top) The domain structure of CHORD protein. The 1 bp deletion caused a frame shift that resulted in a premature stop codon (M172 STOP) in the 2nd CHORD domain. (B–D) Representative images of MARCM clones of the wild-type neuron (B), the CHORD2 mutant neuron (C) and the CHORD2 mutant neuron to which a genomic fragment including the wild-type CHORD gene was introduced (D). The CHORD2 mutation is early larval lethal, and note that all of the data throughout this study are from single neurons homozygous for individual mutations in otherwise heterozygous animals, using the MARCM system (hence, a mosaic analysis), except for the data from RNAi. Scale bars, 50 μm. (E–K) Quantification of branching patterns of dendritic arbors. (E) Dendritic arbor size. (F) Total length of dendritic branches. (G) The number of endings of dendritic branches. (H and I) The branch density: total length/arbor size (H) and ending number/arbor size (I). (J and K) The number (J) and length (K) of branches of each order. All data are presented as means ± standard deviation (SD). *p < 0.05, **p < 0.01, and ***p < 0.001. Blue asterisks indicate statistically significant differences of the cohort from the wild-type neuron; and orange asterisks indicate statistically significant differences of the cohort from the CHORD mutant neuron. NS: Statistically not significant (P > 0.05). (L) Schematic representation of two partly distinct ways of downsizing dendritic arbors. In neurons with defective IIS/TORC1 signaling pathways showed the final arbor pattern similar to the proximal arbor of the normal neuron (left). In contrast CHORD mutant neurons proportionally downsize the original structure, making a miniature form (right).
Whole-genomic sequencing and complementation mapping using Drosophila deficiency stocks identified a 1 bp deletion that leads to a frameshift in the CHORD/morgana gene (denoted CHORD hereafter; Figure 2A). CHORD is an evolutionarily conserved co-chaperone of HSP90, and it negatively regulates Rok kinase (Rok) activity to suppress overduplication of centrosomes. Introduction of the 4757-base-pair (bp) genomic fragment including the CHORD gene almost fully rescued the above phenotypes to normal, demonstrating that CHORD is the causative gene for the miniature dendrite phenotype (Figure 2D and 2E–2K).

The miniature arbor of the CHORD mutant neuron may mimic a default state of the wild-type neuron. One possible explanation of the CHORD phenotype is that the mutant neuron no longer interprets favorable or unfavorable extracellular conditions (either the nutrition and/or body size) and generates the miniature arbor as an invariable default. Alternatively, the CHORD mutant neuron misreads the size of the abdominal lateral plate; consequently, the arbor occupies only a portion (e.g., 50% instead of 100%) of the body surface. To distinguish these possibilities, we examined dendritic arbors of CHORD mutant neurons under the starved condition (Figure 3A and 3B). Neither the size nor the branching complexity significantly changed between the two nutrition conditions (Figure 3C–3I), showing that the CHORD mutant neuron formed the miniature arbor irrespective of the extracellular condition. This result suggested that the latter possibility was less likely. It appears that the wild-type neuron possesses a CHORD-dependent mechanism that extends the branch segment length beyond a preset value in response to a favorable environment.

Elongation/retraction rate of terminal branches is critical for reproducing the miniature phenotype of the CHORD mutant neuron. We next investigated the dynamics of dendrite formation whereby the CHORD mutant neuron produced the miniature arbor during pupal stages. First we found that the mutant arbor underwent a persistent increase in complexity and size, but the overall appearance had already become miniature-like before eclosion (Supplementary Figure S2A–S2H). Then we performed time-lapse recordings of the wild-type and mutant neurons at 70–75 hr after puparium formation (APF), when dendritic arbors were under active construction, and dissected the dendrite dynamics of elongation/retraction and branching.

We quantified frequencies of elongations and retractions of terminal branches (branches of Strahler order 1, see Figure 1L) and found that these parameters were not altered in CHORD mutant neurons (Supplementary Figure S2I and S2J). We also used a Fano Factor as a quantitative descriptor of branch tip dynamics (see details in Methods) and observed that there was no significant difference between the wild-type and CHORD mutant neurons (Supplementary Figure S2K). In contrast, the rates of branch–length changes (µm/min) were significantly decreased in the mutant neurons (Supplementary Figure S2L). In this sensory neuron, most of the branches are collaterals that sprout from stalks, rather than bifurcations of branch tips. Therefore, we quantified the frequency of lateral branching and found that wild-type and CHORD mutant neurons showed similar values (Supplementary Figure S2M). Altogether, dendritic dynamics of CHORD mutant neurons are very similar to that of wild-type neurons, with the exception that the rates of change of branch lengths decrease in the mutant neurons.

We then asked whether these changes can explain the miniature phenotype of the mutant neurons or not. We addressed this question with a computer-assisted simulation of dendrite growth using experimental values. The elongation and retraction of the terminals was represented as the addition and removal, respectively, of one unit segment per 1 minute (=1 step in simulation) at the terminal of the dendrites, and lateral branching was reproduced by the addition of unit segments to the existing branches. The probabilities of elongation, retraction, and lateral branching were determined by experimental data (see details in Methods and Supplementary Table S3). We defined the unit length of a segment as 0.9 µm or 0.75 µm, based on the experimental data for the wild-type or CHORD mutant neurons, respectively (Supplementary Figure S2L). Representative images of the wild-type and mutant model neurons closely resembled those of in vivo neurons (Supplementary Figure S2N). In fact, our quantification showed decreases in the arbor size and the total length, an unaltered ending number,
and an increase in the branch density of the mutant model arbors, compared to the wild-type model arbors (Supplementary Figure S2O–S2S). Thus, these results support the hypothesis that subtle but significant differences in the rates of branch-length changes play a critical role in scaling the dendritic arbor, and that the miniature arbor of the CHORD mutant neuron can be attributed to the scaled-down elongation/retraction dynamics of dendritic branches.

Finally, we examined whether the dendrite dynamics in starved wild-type animals is comparable to that of CHORD mutant neurons. We found that the wild-type neurons under the starved condition showed significantly decreased rates of branch–length changes compared to those under the fed condition, while other parameters were not significantly changed (Supplementary Figure S2I–S2M), very much resembling CHORD mutant neurons under the fed condition. These results strengthened our hypothesis that coordinated regulation of the elongation/retraction dynamics of branches contributes to scaling of the dendritic arbor.

### Genetic interaction between CHORD and the TORC2 component Rictor

We next asked how body size/nutrition state is conveyed to CHORD. As a candidate of such an upstream molecule, we examined the involvement of TORC2, another TOR complex, for the following reasons: (1) TORC2 is activated by association with ribosomes in a growth factor dependent manner, thus it may mediate between the extracellular environment and growth capacity of individual cells. (2) TORC2 regulates the actin cytoskeleton and controls dendrite pattern formation. We assessed the phenotype of mutant neurons of rictor, which encodes an essential and specific component of TORC2 (Figure 4A and 4B). rictor mutant neurons showed complicated but intriguing phenotypes; that is, combined features of undergrowth and miniaturization. They showed a decrease in the arbor size, the total length, and the ending number (Figure 4D–F), which is characteristic of the undergrowth phenotype. On the other hand, they also exhibited features of the miniature phenotype: increases in the branch density (both the total length/arbor size and the ending number/arbor size, as shown in Figure 4G and 4H) and a decrease in segment length (Figure 4I). Importantly, overexpression of the CHORD transgene in rictor mutant neurons partially restored the features of the miniature phenotype (Figure 4G–4I), but not those of the undergrowth phenotype (Figure 4D–4F). In contrast to this genetic interaction between CHORD and TORC2 (Rictor), CHORD overexpression did not rescue simplified phenotype of the dnr knockdown (Supplementary Figure S1M and S1N). All these results could be explained by the hypothesis that CHORD functions downstream of TORC2, being at least partly separate from TORC1. The partially separate nature was further supported by our KD experiment using S2 cells (Supplementary Figure S1O). Even when CHORD was knocked down, phosphorylation of S6K (a readout of TORC1 activity) was up-regulated in an insulin-dependent manner, as in the control knockdown cells.

We then asked whether the TORC2 activity is altered under the nutrient-limited condition that was employed to decrease body size. We starved larvae for 8 hrs beyond 91–96 hr AEL and analyzed the phosphorylation of Akt at Ser505 as an established readout of the TORC2 activity. The level of S505 phosphorylation was significantly and reproducibly reduced under this condition compared to that in well-fed flies (Figure 4K). All these results suggest the possibility that TORC2 may contribute to the extracellular conditions (body size and/or nutrients) to CHORD proteins (Figure 4L).

### Discussion

This study provides novel mechanistic insights into size control of neuronal dendritic arbors. To sample sensory input precisely, certain types of neurons should adjust their dendritic arbor size to cover the receptive field completely. We showed that there are two distinct ways of downsizing dendritic arbors when the field size is reduced: one way is to arrest both growth and branching during maturation of dendrite morphologies, causing the “undergrowth” phenotype. The other way is to regulate the elongation of dendrites (more specifically speaking, the length of branch segments) selectively, thereby making a “miniature” form (Figure 2L). Our results support the notion that dendritic growth and branching are controlled by at least partly separate mechanisms, which is also seen in other instances such as the development of postembryonic dendritic architecture of motor-neurons. At the molecular level, the IIS/TORC1 pathways control both growth and branching to avert underdevelopment, whereas CHORD and TORC2 tunes the segment length to realize proportional scaling of the entire arbor (Figure 4L). Ablation of the TORC2 component Rictor in mouse Purkinje cells causes multiple structural changes of dendritic arbors, including a decrease in total dendrite length and an increase in the number of primary branches. It will require further investigation to address how the overall branching complexity and the segment length is affected in the absence of Rictor, and to fully characterize the loss of function phenotype of CHORD in this subtype of neurons.

CHORD was originally discovered in plants as a key player in the innate immune response, in animals, CHORD negatively regulates Rho kinase activity, thereby suppressing overduplication of centrosomes. CHORD is also expressed in tissues that are populated by postmitotic cells, such as brains, and has been proposed to function beyond regulating cell division. Here, we revealed that CHORD regulates the size of dendritic arbors: cells defective in CHORD showed a decreased elongation/retraction rate of terminal branches; thus, by extrapolation, the normal function of CHORD must be to accelerate the elongation and retraction of terminal branches. Rho-kinase (Rok) is reported to inhibit neurite outgrowth, dendritic branching, and spine formation. Therefore, we pursued the possibility that Rok acts downstream of CHORD. However, we couldn’t find any genetic and biochemical evidence for the interaction between Rok and CHORD (Supplementary Figure S3; see details in the legend). Instead, our results suggested that CHORD is functionally related to the TORC2 signaling pathway, which regulates the actin cytoskeleton. It has been reported in mammalian fibroblasts that phosphorylation by TORC2 facilitates folding of some AGC kinases, such as Akt and conventional PKCs, and stabilizes them, and that newly synthesized, unphosphorylated Akt is protected by HSP90 from degradation. Therefore, CHORD may contribute as a co-chaperone to the stability of the AGC kinases by recruiting HSP90 to those clients, or it may employ its own chaperone activity to perform this task. This hypothetical role of CHORD might underlie the partial rescue of the rictor mutant neuron by CHORD overexpression.

Comparative anatomical studies have reported that dendrites of some types of neurons, such as sympathetic neurons, become larger and increase branching complexity as the brain or the body size increases across species, which we designate overgrowth as opposed to undergrowth. In contrast, other types of neurons such as somatosensory thalamocortical projection neurons increase their dendritic arbor size with larger brains, while preserving key features of the dendritic branching pattern, which is proportional magnification. These two distinct ways of scaling, overgrowth/undergrowth and miniaturization/magnification, might be regulated by IIS/TORC1 and TORC2/CHORD, respectively, and may contribute to neuronal cell-type specific information processing.

It should be noted that “scaling growth” of dendritic arbors does take place during larval development, but mechanistically it is distinct from the arbor scaling at postlarval stages, which we reported in this study. Larval da neurons regulate their growth in coordination with the expanding body whose mass increases by approximately 200-fold during the complete larval development. This is
accomplished by increasing both branching numbers and the total length, where IIS/TORC1 plays a critical role \(^{29,45}\), but CHORD appears to be dispensable (Supplementary Figure S2), and TORC2 is required for dendritic tiling for larval da neurons\(^{29}\). On the other hand, the da neuron during pupal development first prunes its dendrites and then starts constructing arbors for adult life; thus this neuron completes this task in a body whose volume has been predetermined by the nutritional status during larval development. It is this task that requires the role of CHORD, which fits the dendritic arbor to the final adult body size.

We speculate that this hypothetical CHORD-dependent mechanism somehow senses body size by receiving as-yet undetermined signals, either local or systemic, from other tissues. Candidate tissues for sources of such signals would be either those adjacent to the neuron, such as lateral tergosternal muscles, abdominal histoblasts (epithelial cells), and glial cells, and/or tissues that secrete growth factors, such as fat body and insulin producing cells\(^{8,45–48}\). Therefore the dendrite scaling of the adult da neuron might provide a useful model system to study interactions between neurons and other organs.

**Methods**

**Molecular cloning.** Six tandem repeats of a 20-base-pair (bp) sequence that includes a proneural binding site ([scE1]\(^{6}\))\(^{49}\) were used to build a SOP-FLP transgene. The hsp70 minimal promoter and the entire coding sequence of flp from the UAS-FLP vector (DGRC) were inserted between [scE1]\(^{6}\) and a SV40-polyA sequence. This SOP-FLP transgene was cloned into the pHStinger vector, or a pUAST vector from which the UAS sequence and hsp70 minimal promoter had been removed. To generate a rescue construct, the 4754-bp sequence (3R: 20009616-20013777 in version FB2013_05) that includes CHORD was amplified from yw genomic DNA and cloned into the vector pCasper. To generate UAS shRNA for each gene, we followed a protocol previously described\(^{50}\). The target sequences of the shRNAs are as follows:

\[
\text{5’-CACCGAGTTCCTCAACATCAA-3’}
\]

\[
\text{5’-TTCGACCTGGATGACATTA-3’}
\]

These shRNAs were cloned into the vector pUAST-attB. All constructs were injected, in accordance with standard protocols, to generate transgenic fly lines.

**Drosophila strains.** Detailed genotypes of the animals and clones are described and summarized in Supplementary Table S1. To visualize dendrites and/or express...
transgene, we used the following Gal4 drivers: Gr28b-Gal4 and Gal80-Gal4. To express fluorescent proteins, we used UAS-mCD8::GFP (#5137 of the Bloomington Stock Center) or UAS-Venus-pm::GFP (#4350. Other strains used were dpy11; Akt[17], Tor[5–7], rok[3–14], UAS-disc2 (#60009 of Vienna Drosophila RNAi Center), UAS-raptor[34] (HMS0124/83418 of the Bloomington Stock Center), UAS-CherryRV[35] (335785 of the Bloomington Stock Center), UAS-CHORDRNAi[36] (this study), UAS-Dp110[9841A] (#25910, Bloomington Stock Center), UAS-rok[36] (GD1522/403 and KK107802/#104675 of the Vienna Drosophila RNAi Center; FG3255/82797, HMS01311/#34324, and GL00209/#35305 of the Bloomington Stock Center), UAS-lsp[90] (HMS08989/#33947 of the Bloomington Stock Center) and UAS-Rok.CAT[17].

MARCM-based forward genetic screen. The piggyBac insertion collection with FRT insertion[37] was used for genetic screening. We crossed individual insertion stocks to "SOP-FLP based MARCM-ready" fly stocks for (2nd chromosome left arm screen, Gal4+>UAS-Venus-pm SOP-FLP[20]; tubP-Gal80 FRT40A; for 3rd chromosome left arm screen, Gal80-Gal4 UAS-FLP[20]; with UAS-scrib[20]; FRT40A; and for 3rd chromosome right arm screen, hsh2P UAS-mCD8::GFP, Gal4+>UAS-mCD8::GFP SOP-FLP[20]/CyO; FRT82b tubP-Gal80). We then mounted adult abdomens, and dendrites were imaged under a laser scanning microscope. We screened 1537 piggyBac stocks and isolated 3 stocks (L04611, L04133, and L03277) that showed the "miniature" phenotype, and 19 that showed the "undergrowth" phenotype (Supplementary Table S2). Both L04611 and L04133 were homoyzogous lethal.

Mapping and identification of the mutation that was responsible for the "miniature" phenotype. Starting from the above L04611 and L04133 stocks of the piggyBac collection, we generated precise excision strains. These precise excision strains were still homoyzogous lethal, and MARCM clones of these strains showed the miniature dendritic phenotypes. We performed a complementation test for lethality between L04611 and L04133, and found that they belong to the same complementation group. These results suggested that the lethality and the "miniature" dendrite phenotype were caused by a background mutation(s) that was not linked to the piggyBac insertion. We backcrossed the excision strains to the control FRT82b stock three times and used the progeny for the phenotypic analysis below.

To identify the mutated gene that gave rise to the phenotype, we determined whole-genomic sequences of the two mutant strains with an Illumina next-generation sequencer and directly compared them with each other. For the sequencing, we prepared genomic DNA from larvae that were heterozygous for L04611 or L04133 and employed paired-end Illumina sequencing technology. We also determined the sequences of L02779 and L00232, whose MARCM clones did not show a miniature phenotype and negative controls. Then we mapped the sequence data to the wild-type reference genome and identified variants relative to the wild-type reference genome by using Burrows-Wheeler Aligner (BWA) and SAMTools. In addition to the whole genome sequencing, we performed conventional deficiency mapping of the lethality by using the Bloomington Deficiency Kit, and mapped the mutation responsible for the lethality (and possibly the clone phenotype) within a ~45 kb region in the right arm of the 3rd chromosome. Within this ~45 kb mapped region, we found only one mutation unique to L04611 and L04133: a 1 bp deletion in the CHORD/morgana gene that caused a frame shift (denoted as CHORD), which was confirmed by the Sanger method.

Image collection and image analysis. Imaging da neurons in whole-mount adults was done as described earlier[18]. Briefly, we collected adult females within 12-hour after eclosion unless described otherwise, washed them in 0.7% NaCl and 0.3% Triton X-100, removed the heads and legs of adult flies, and mounted the abdomens in 50% glycerol on slides, between spacers made of vinyl tape. All of the images were acquired using a Nikon C1 laser-scanning confocal microscope or a Zeiss LSM 510 META laser-scanning confocal microscope.

For quantification of dendritic patterns of da neurons, live imaged dendrites were used for measurement, each pupa was kept at 25°C under normal light conditions. For quantification, we folded legs and put abdomens on the dish and tilted them to retain an imaging session, divided by the mean length of that branch[35]. We quantified neurons whose FF was above 0.35, which is defined as dynamic neurons[36].

Reconstruction of dendrite dynamics in a computer-aided simulation. Dendrite growth dynamics was modeled in two-dimensional space following previous mathematical models[38,39]. The 'elongation' and 'retraction' were represented by the addition and removal of 1 unit segment, respectively. As a length of 1 unit segment, we employed 0.9 μm and 0.75 μm to recapitulate the wild-type and a CHORD mutant neurons, respectively. Lateral branching was reproduced by the addition of unit segments to the existed branches. The angle of elongation was assumed to follow a homogeneous distribution from −7.5° to +7.5°, while that of branching was assumed to follow a normal distribution with average 97° and standard deviation 38°. The initial condition was as follows: 2 branches with 20-segment lengths sprout from the origin in anti-parallel directions along the y axis and bifurcate with angle 30°, and each of these 4 branches elongate 10-segments in length. Other parameters used in the simulation are summarized in Supplementary Table S3.

Production of antibodies, RNAi, and Western blotting. Guinea pigs were immunized with GST proteins that had been fused to the N-terminal 200 amino acids of CHORD to generate anti-CHORD antibody. For RNAi experiments, S2 cells were cultured with 20 μg/ml dsRNA of the full-length CHORD or GFP-coding sequence for 3 days. Lysates were separated on a 8% polyacrylamide gel and transferred to PVDF (Millipore). Antibodies were used at the following concentrations: guinea pig anti-CHORD, 1: 1000 (this study); rabbit anti-phospho-Mycosin light chain 2 (Ser-19, 1: 1000 (Cell Signaling #3671); rabbit anti-phospho-Akt (Ser-505), 1: 1000 (Cell Signaling #4054); rabbit anti-Akt, 1: 1000 (Cell Signaling #9227); mouse anti-actin, 1: 1000 (Millipore MAB1051). Signals were detected with ECL plus (Amersham). To obtain stronger signals, we employed Can Get Signal Immunoreaction Enhancer solution (TOYOBO).


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