Glia Promote Local Synaptogenesis Through UNC-6 (Netrin) Signaling in *Caenorhabditis elegans*

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Neural circuits are assembled through the coordinated innervation of pre- and postsynaptic partners. We show that connectivity between two interneurons, AIY and RIA, in *Caenorhabditis elegans* is orchestrated by a pair of glial cells that express UNC-6 (netrin). In the postsynaptic neuron RIA, the netrin receptor UNC-40 (DCC, deleted in colorectal cancer) plays a conventional guidance role, directing outgrowth of the RIA process ventrally toward the glia. In the presynaptic neuron AIY, UNC-40 (DCC) plays an unexpected and previously uncharacterized role: It cell-autonomously promotes assembly of presynaptic terminals in the immediate vicinity of the glial cell endfoot. These results indicate that netrin can be used both for guidance and local synaptogenesis and suggest that glial cells can function as guideposts during the assembly of neural circuits in vivo.

Neural circuit formation requires an intricate orchestration of multiple developmental events, including cell migration, axon guidance, dendritic growth, synaptic target selection, and synaptogenesis (1–3). These developmental events are coordinated in pre- and postsynaptic neuronal partners to form the functional neural circuits that underlie behaviors. Although the organization and specificity of these neural circuits is well documented, the cellular and molecular mechanisms that underlie their precise development are not well understood.

To explore how precise neural connectivity is achieved, we studied the synaptic connections between two interneurons in the *C. elegans* brain: presynaptic AIY and postsynaptic RIA. These two interneurons navigate complex cellular environments, discriminating among multiple potential targets before finding and innervating each other at a discrete region of their respective processes (4). We generated single-cell fluorescent markers to visualize AIY-RIA connectivity in vivo and observed a discrete clustering of presynaptic AIY markers in a segment of the process we termed zone 2. This zone appears to be the specialized presynaptic region where AIY forms synapses onto RIA, as well as RIB and AIZ neurons. First, the fluorescently labeled presynaptic protein Rab-3, ELKS-1, and SYD-2 are all more concentrated in zone 2 than in other regions of the axon (Figs. 1A and 2B and fig. S4A). Second, these markers cluster at the exact location at which AIY to RIA synapses are seen in electron micrographs of wild-type animals (fig. S1M) (5). Third, this region has a wider diameter than other regions of the axon, a property that we found to be uniquely associated with the presynaptic region of AIY in electron micrographs (fig. S1, A and M to Q). These combined properties were taken as evidence of presynaptic differentiation and were very reproducible across animals (Fig. 1 and fig. S1).

Reconstructions of electron microscopy (EM) micrographs (5) revealed that AIY has three distinct anatomical regions throughout its process: a segment proximal to the AIY cell body that is devoid of synapses (zone 1); the synapse-rich region where AIY forms synapses onto RIA, AIZ, and RIB just as the AIY process turns dorsally (zone 2); and a distal axon segment within the nerve ring that has four to eight small presynaptic specializations (zone 3).

To identify the molecular signals that direct this precise innervation, we performed a visual genetic screen for mutants with an abnormal synapse distribution in AIY. From this screen, we isolated the *wy871* mutation, an allele of *unc-40* (fig. S2). *Unc-40* (DCC, deleted in colorectal cancer) is a transmembrane immunoglobulin superfamily protein that is a receptor for the axon guidance molecule UNC-6 (netrin) (6, 7). *unc-40* animals had no detectable axon guidance defects in AIY except for an axon truncation defect observed in 7.8% of the animals (n = 153 animals; fig. S3). However, they showed a highly penetrant defect in the presynaptic specialization of AIY at zone 2: 95.3% of *unc-40* (*wy871*) animals displayed a severe reduction of active zone markers ELKS-1::YFP (yellow fluorescent protein) and SYD-2::GFP (green fluorescent protein) and a synaptic vesicle marker, mCherry::RAB-3, in zone 2 (n = 128 animals; Fig. 2, A to K, and fig. S4). In addition, the AIY axon diameter in zone 2 failed to widen into the characteristic presynaptic varicosity seen in wild-type animals (fig. S5). By contrast, in the more-dorsal zone 3 synaptic regions, *unc-40* animals had normal or increased levels of synaptic vesicle proteins and a normal or increased diameter (Fig. 2, F to I, and fig. S5). These defects suggest a specific defect in the presynaptic differentiation of AIY in zone 2, although a detailed analysis of AIY synaptic ultrastructure and function could...
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reveal abnormalities in other AIY synapses. Although unc-40 animals do not have substantial AIY axon guidance defects, RIA axon guidance is severely affected in unc-40 mutants: RIA processes fail to extend ventrally to create the loop that is innervated by AIY in zone 2 (95.8% penetrant, \( n = 191 \) animals; fig. S3).

To identify the cell(s) in which unc-40 functions to direct AIY-RIA innervation, we analyzed unc-40 mosaic animals that retain an unstable rescuing array in subsets of cells (8). Interestingly, only when the array was retained in the AIY interneuron did we observe significant rescue of the AIY presynaptic patterning defects (\( P < 0.001; \) Fig. 2, L to M, and figs. S5 and S6). Retention of the array in the closely related interneuron RIB or RIM (RIB/RIM) (9) did not result in rescue. Retention in the RIA interneuron resulted in rescue of the RIA axon guidance defect but not of the AIY presynaptic phenotype (fig. S7).

Together, these data indicate that axon guidance of RIA and patterning of presynaptic specializations in AIY are independent, cell-autonomous events. These mosaic analyses are also consistent with the observation that UNC-40 (DCC) is endogenously expressed in AIY interneurons (fig. S6) and indicate a previously uncharacterized role for UNC-40 (DCC) in interneurons (fig. S6) and indicate a previously uncharacterized role for UNC-40 (DCC) in interneurons (fig. 3, A and C; 71.5% animals with enriched localization, \( n = 214 \) animals). This enrichment is dependent on UNC-6 (netrin), because UNC-40 (DCC) failed to concentrate at the presynaptic sites of AIY in unc-6 animals (Fig. 3, B and C; 87.6% with diffuse localization, \( n = 201 \) animals). Consistent with this observation, in unc-6 animals circuit assembly between AIY and RIA is abnormal, phenocopying the unc-40 mutant defect (fig. S8).

During neurulation, when AIY-RIA innervation takes place, UNC-6 (netrin) is exclusively expressed by ventral cephalic sheath cells (VCSCs) at the nerve ring (10). VCSCs are non-neuronal cells that are morphologically similar to vertebrate astrocytes (11). Mosaic analysis using an unc-6 rescuing construct showed that expression of UNC-6 (netrin) by VCSCs regulates the pattern of AIY presynapses (fig. S8B, \( P < 0.001 \)).

VCSCs have slender processes that enter the nerve ring and terminate at specific synaptic sites, one of them being zone 2, the region where AIY innervates RIA (4). We investigated the physical relations between the VCSCs, AIY, and RIA and observed that the VCSCs project endfeet to the anterior end of the ventral nerve cord and form deeply invaginated membrane lamellae that ensheath the zone 1 region where AIY innervates RIA. The relative positioning of the VCSC endfeet with respect to AIY and RIA is extremely stereotyped across animals, revealing a tight and reproducible anatomical relation between the UNC-6–secreting VCSCs and the AIY-RIA synapses (Fig. 3, D to J, and fig. S1).

To test further whether the precise anatomical relation between the sheath cells and AIY-RIA synapses is instructive in mediating AIY-RIA innervation, we identified mutants that disrupt sheath cell morphology. We found that mutations in unc-34/enabled, a regulator of the actin cytoskeleton, altered the morphology of cephalic sheath cells: 47.8% of the unc-34 animals have distended endfeet, which project further posteriorly \( (n = 94 \) animals; Fig. 4, A and D, and fig. S9A). In these unc-34 animals, the distended VCSCs ectopically ensheath zone 1 of AIY in addition to zones 2 and 3 (Fig. 4, A and D). Also in these animals, we observed a change in the distribution of UNC-40 (DCC), which ectopically localized to zone 1 (Fig. 4, E to G). Furthermore, there was a concomitant change in the distribution of AIY presynapses: ectopic AIY presynaptic specializations now formed in zone 1, the region of overlap with distended sheath cell endfeet where ectopic UNC-40 localized (Fig. 4, B and D). Moreover, RIA axon guidance changed accordingly, with the RIA postsynaptic loop extending further posteriorly to the region covered by the sheath cell endfeet \( (P < 0.001; \) Fig. 4, C and D, and fig. S9A).

An epistasis analysis confirmed that the displacement of presynaptic sites observed in unc-34 animals is an UNC-40 (DCC)–dependent event \( (n = 94 \) animals circuit assembly between AIY and RIA. The glial cells mark a neuospatial coordinate in the C. elegans nerve ring through the secretion of UNC-6 (netrin). This in turn signals to AIY and RIA neurons through UNC-40 (DCC). UNC-40 (DCC) simultaneously activates two different and independent pathways in each neuron, orchestrating presynaptic assembly in AIY and axon guidance of postsynaptic RIA to the glial-specified location.

How can the same receptor and ligand elicit diverse cellular responses in distinct neurons? UNC-40 (DCC) has been reported to regulate diverse developmental processes like cell migration, neuronal polarization, and axon guidance, all of which involve an initial polarization event that may be mediated by netrin signaling \((6, 7, 12–15)\). The patterning of AIY presynaptic regions can also be considered a local polarization event, through which AIY transforms a region of its plasma membrane into a specialized presynaptic area. Other guidance molecules, such as the Eph family of receptors and their ephrin ligands, have been shown to play roles in growth cone guidance as well as the development of mature excitatory synapses \((16)\). In ephrin-mediated signaling, distinct cellular responses are likely generated by the developmental context and by diverse downstream targets \((17)\). Similar mechanisms could explain the distinct cellular responses to UNC-40 (DCC) signaling.
Fig. 2. UNC-40 (DCC) is required for correct presynaptic patterning in AIY. (A to J) Distribution of presynaptic sites of AIY in wild-type [(A) to (E)] or unc-40 [(F) to (J)] animals. Confocal micrographs demonstrate the colocalization and patterning of AIY presynaptic vesicles [mCherry::RAB-3 pseudocolored red in (A) and (F)] and active zones [ELKS-1::ERC::YFP pseudocolored green in (B) and (G)] in a representative wild-type [(A) to (C)] or unc-40 [(F) to (H)] animal, evident in the double-label merge images [(C) and (H)] and represented in the diagram of presynaptic site distribution [(D) and (I)]. Three-dimension line scan profiles of the fluorescence intensity (arbitrary units, AU) of UNC-40 in AIY and RIA (L) or RIB/RIM (M) were scored for retention of the UNC-40::GFP in AIY of a representative wild-type (A) or unc-6 mutants (B). (C) Quantification comparing the enrichment of UNC-40::GFP in the presynaptic regions of wild-type versus unc-6 animals. Two different transgenic lines (lines 1 or 2), both expressing UNC-40::GFP, were scored. (D) Projection of confocal micrographs obtained from a representative animal simultaneously expressing hib-17::mCherry (to label ventral cephalic sheath cells and pseudocolored blue), ttx-3::eGFP::rab-3 (to label presynapses in AIY and pseudocolored red), and glr-3::glr-1::YFP (to label the glutamate receptors in RIA and pseudocolored green). (E) Volume rendering of (D) showing the relative positioning of the glial-like ventral cephalic sheath cells with respect to the region of innervation between AIY and RIA (29). (F) Diagram of (D) and (E). The asterisk marks the posterior extension of the sheath cell. (G) Volume rendering of the ventral sheath cell in (D). Note the groove in the boxed region where AIY and RIA are ensheathed by the ventral cephalic sheath cell and innervate each other. (H and I) Single confocal plane of (D), with presynaptic vesicles in AIY [labeled with cyan fluorescent protein (CFP)::RAB-3 and pseudocolored red in (H)], glutamate receptors in RIA [labeled with GLR-1::YFP and pseudocolored green in (I)] and a triple label including expression of the mCherry cytoplasmic fluorophore in the sheath cells [pseudocolored blue in (J)]. This anatomical relation was extremely stereotyped across animals (n = 500). Scale bars, 5 μm.

Fig. 3. Ventral cephalic sheath cells control UNC-40 (DCC) enrichment in presynaptic regions by secreting UNC-6 (netrin). (A and B) Localization of UNC-40::GFP in AIY of a representative wild-type (A) or unc-6 (B) animal. Note UNC-40::GFP enrichment in zones 2 and 3 in wild-type animals [(A) dashed box], as compared to diffuse localization in unc-6 animals (B). (C) Quantification of the enrichment of UNC-40::GFP in the presynaptic regions of wild-type versus unc-6 animals. Two different transgenic lines (lines 1 or 2), both expressing UNC-40::GFP, were scored. (D) Projection of confocal micrographs obtained from a representative animal simultaneously expressing hib-17::mCherry (to label ventral cephalic sheath cells and pseudocolored blue), ttx-3::eGFP::rab-3 (to label presynapses in AIY and pseudocolored red), and glr-3::glr-1::YFP (to label the glutamate receptors in RIA and pseudocolored green). (E) Volume rendering of (D) showing the relative positioning of the glial-like ventral cephalic sheath cells with respect to the region of innervation between AIY and RIA (29). (F) Diagram of (D) and (E). The asterisk marks the posterior extension of the sheath cell. (G) Volume rendering of the ventral sheath cell in (D). Note the groove in the boxed region where AIY and RIA are ensheathed by the ventral cephalic sheath cells and innervate each other. (H and I) Single confocal plane of (D), with presynaptic vesicles in AIY [labeled with cyan fluorescent protein (CFP)::RAB-3 and pseudocolored red in (H)], glutamate receptors in RIA [labeled with GLR-1::YFP and pseudocolored green in (I)] and a triple label including expression of the mCherry cytoplasmic fluorophore in the sheath cells [pseudocolored blue in (J)]. This anatomical relation was extremely stereotyped across animals (n = 500). Scale bars, 5 μm.
Although cell-specific events in AIY and RIA happen independently of each other, their simultaneous regulation by a single molecule allows coordinated circuit assembly. UNC-6 (netrin) is a secreted chemotrophic factor that can act as a long-range chemical cue or a short-range signaling molecule, depending on the developmental context (18–20). Given the close anatomical relation between the source of UNC-6 (netrin) (VCSCs) and the AIY:RIA synapses, it is likely that UNC-6 (netrin) acts as a short-range signaling molecule in this pathway, specifying microenvironments that promote UNC-40 (DCC) enrichment and synaptogenesis. Indeed, vertebrate astrocytes have been shown to be functionally compartmentalized into subcellular microdomains, which may be important in regulating localized secretion of signaling molecules regulating synaptic assembly and function (21). Furthermore, UNC-6 (netrin) has been reported to mediate adhesive interactions and function as a short-range target recognition molecule in other developmental events (18, 19, 22).

Our study is consistent with observations made in vertebrates and highlights the importance of glial cells in specifying precise neural connectivity. The fact that dissociated cultured neurons can form functional synapses in the absence of glial cells suggests that pre- and postsynaptic neurons are sufficient to assemble chemical synapses. However, growing evidence suggests that glia are essential regulators of synaptic assembly and function in vivo (11, 23–26). For instance, astrocyte-secreted thrombospondin increases the density of synapses in the mammalian central nervous system (23, 27).

Our data show that glia can also specify neural connectivity in vivo by marking a neurosponal coordinate, which achieves precise circuit assembly by controlling both synaptic partner choices and the location of innervation. Such stereotypic synaptic assembly gives rise to highly organized neuropil structures such as the nematode nerve ring. Similarly organized neuropil structures are also evident throughout the vertebrate central nervous system, exemplified by the stratified organization of the inner plexiform layer of the vertebrate retina and the glomeruli in the olfactory bulb (2, 28). The roles observed here for glia in C. elegans may be evolutionarily conserved, such that proteins with multifunctional roles in spatial patterning, like UNC-40 (DCC), would detect the localized expression of gli signals and thus orchestrate the formation of neural circuits.

References and Notes

9. Materials and methods are available on Science Online.
27. K. S. Christopherson et al., Cell 120, 421 (2005).
29. We thank G. Wang, C. Johnson, J. Aduhya, M. Nonet, Y. Kohara, J. Kaplan, G. Garriga, E. Lundquist, the Caoorhabditis Genetic Center, and the Japanese National BioResource Project for strains and reagents; D. Hall and Z. Altun for diagrams used in the figures; C. Bargmann for helpful discussions and generous sharing of advice and reagents; C. Gao and Frank Chen for technical assistance; and S. Margolis, L. Looger, J. Irazoqui, B. Barres, and members of the Shen lab for thoughtful comments on the manuscript. This work was funded by grants to K.S. from the following: the McKnight Endowment Fund, the W. M. Keck Foundation, and the Searle Scholar program. M.A.M. was supported by the Stanford Medical Scientist Training Program and NIH grant GM007365. D.A.C.-R. was supported by the Damon Runyon Foundation and NIH grant K99 NS057931-01.

Supporting Online Material

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Materials and Methods
Figs. S1 to S9

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