sli is required for proper morphology and migration of sensory neurons in the Drosophila PNS

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Abstract

Neurons and glial cells coordinate with each other in many different aspects of nervous system development. Both types of cells are receiving multiple guidance cues to guide the neurons and glial cells to their proper final position. The lateral chordotonal organs (lch5) of the Drosophila peripheral nervous system (PNS) are composed of five sensory neurons surrounded by four different glial cells, scolopale cells, cap cells, attachment cells and ligament cells. During embryogenesis, the lch5 neurons go through a rotation and ventral migration to reach their final position in the lateral region of the abdomen. We show here that the extracellular ligand sli is required for the proper ventral migration and morphology of the lch5 neurons. We further show that mutations in the Sli receptors Robo and Robo2 also display similar defects as loss of sli, suggesting a role for Slit-Robo signaling in lch5 migration and positioning. Additionally, we demonstrate that the scolopale, cap and attachment cells follow the mis-migrated lch5 neurons in sli mutants, while the ventral stretching of the ligament cells seems to be independent of the lch5 neurons. This study sheds light on the role of Slit-Robo signaling in sensory neuron development.

Keywords: Sli, Chordotonal neurons, PNS, Drosophila

Introduction

The nervous system is made up of neurons and glial cells. Proper development of the nervous system requires coordination between these two cell types. It is known that glial cells ensheath axons and function to drive nerve formation, provide trophic support for neuronal survival, provide guidance during axon pathfinding, and modulate dendrite morphology [1–4]. Even with all this information on the multitude of glial cell functions, we still have much to learn about how specific glial cells coordinate with neurons in the formation of the nervous system.

The fruit fly Drosophila melanogaster is a well-known model system for studying many of the fundamental aspects of neural development, including neuron-glia interactions [1–4], and the mechanisms and signaling pathways necessary for axon guidance [5–7]. For example, the Slit-Robo signaling pathways was found to be required for proper crossing of commissural axons in the central nervous system (CNS) in Drosophila [8–12]. In this study, we are examining more closely the role of slit (sli), and possible Slit-Robo signaling in the embryonic peripheral nervous system (PNS).

The embryonic PNS of Drosophila melanogaster is composed of different types of sensory neurons, which are divided into Type I - neurons with single dendrites, and Type II - multi-dendritic neurons. Type I neurons are divided further into four clusters, dorsal (d), lateral (l), ventral’ (v’), and ventral (v), according to their final position along the dorsal-ventral axis of the embryo. The lateral chordotonal (lch5) neurons are a group of five Type I mechanosensory neurons that sense stretch and vibration [13–16]. There is one group of lch5 neurons in each of seven abdominal segments of the Drosophila embryo [14, 17]. The precursors of the lch5 neurons initiate in a dorsal position in the embryo at stage 12 and migrate ventrally to their final lateral position at stage 15 (Fig. 1A, B) [13, 15, 17–19]. By stage 15 these neurons have a very distinctive morphology, which includes neuron shape, direction of dendrites and spacing of individual neurons relative to each other (Fig. 1A, B, red cells). The shape of each of the five neurons in the lch5 cluster has a teardrop outline with the single dendrite...
pointing in a dorsal-posterior direction (Fig. 1A, B, red cells). These chordotonal neurons are surrounded by four groups of secondary (glial) support cells, scolopale cells, cap cells and attachment cells that are dorsal to the neurons, and ligament cells that are ventral to the neurons (Fig. 1B, C) [18, 20–22]. The lch5 neurons plus their support cells coalesce into one lateral chordotonal organ (lch5 organ). On the dorsal side, the cap cells are connected to the ectoderm by attachment cells [18, 22]. The scolopale cells surround the tip of the dendrite which may interact with migratory cues along the pathway [17, 20]. On the ventral side, the ligament cells stretch ventrally to attach the lch5 organ to the ectoderm (Fig. 1C) [17, 18, 22].

Unlike most of the neurons in the PNS, the lch5 neurons go through a rotation and migration during embryogenesis. Rotation and migration both occur after stage 12 with rotation followed by migration [17, 18]. Prior to stage 12, the dendrites of all thoracic and abdominal chordotonal neurons, including the lch5 neurons, face ventrally. After stage 12, the abdominal chordotonal neurons rotate until the dendrites face dorsal posteriorly [17, 18]. Although the exact mechanism of this rotation and migration is not known, a few different mutations have displayed lch5 migration and rotation defects. For example, it has been shown that Slit-Robo signaling does affect these two processes. Specifically, it was mentioned, but never shown, that in the absence of the extracellular ligand sli, the cell bodies of lch5 neurons either stay dorsally located or their dendrites are aberrantly oriented. The same defect is observed in a double mutant of the sli receptors robo and robo2 [19].

Likewise, the Robo receptor is expressed at the tips of lch5 dendrites while Robo2 is expressed along the entire lch5 dendrites in abdominal segments [17]. In the thoracic region, Robo2 receptor expressed in the visceral mesoderm binds Slit and presents Slit to Robo receptors expressed on thoracic chordotonal neurons, thereby preventing migration of the thoracic chordotonal organs [17]. Additionally, loss of the transcription factor ventral veinless (vvl) results in a failure of the lch5 chordotonal neurons to migrate ventrally, similar to that seen in the absence of sli [18, 23]. The question we are asking is what role does Slit-Robo signaling play in lch5 neuronal migration and morphology.

In this study, we have sought to shed further light on the role of sli on migration and final morphology of the lch5 cluster. We show here that sli is necessary for the ventral migration, final positioning, and morphology of the lch5 chordotonal neurons in the Drosophila PNS. In addition, we show that the absence of either robo or robo2 display similar, albeit less severe, defects. Further, we show that in the absence of sli, the scolopale cells, cap cells and attachment cells follow the mis- or non-migrated neurons, whereas the ligament cells continue stretching ventrally in the absence of sli. Thus, we have shown that Slit-Robo signaling is important for yet another piece of neural development.

Materials and methods
Drosophila strains
The following Drosophila strains were used in this study: Canton S (CS) as the wild-type (WT) control, sli<sup>2</sup> (sli null
ated anti-mouse and biotinylated anti-rabbit.

555, anti-Rabbit 488, anti-Rat 488, anti-Rat 647, biotinyl-

secondary antibodies were used at 1:500: anti-Mouse

GFP (1:5000, ThermoFisher Scientific). The following

α

anti-Robo (13C9) (1:10, DSHB), rabbit anti-

anti-Repo (1:100, DSHB), mouse anti-Prospero (1:10,

Molecular Probes), mouse anti-22C10 (1:10, Develop-

β

mental Studies Hybridoma Bank (DSHB), University of

Iowa, IA), rat anti-Elav (7E8A10) (1:10, DSHB), mouse

anti-Repo (1:100, DSHB), mouse anti-Prospero (1:10,

DSHB), mouse anti-Slit (C555.6D) (1:10, DSHB), mouse

anti-Robo (13C9) (1:10, DSHB), rabbit anti-α-tubulin

85E (1:50, Thermo Fisher Scientific), and rabbit anti-

GFP (1:5000, ThermoFisher Scientific). The following

secondary antibodies were used at 1:500: anti-Mouse

555, anti-Rabbit 488, anti-Rat 488, anti-Rat 647, biotinyl-

ated anti-mouse and biotinylated anti-rabbit.

Staining procedures

HRP and fluorescence immunohistochemistry were per-

formed as described previously [26–28]. The following

primary antibodies were used: rabbit anti-β-gal (1:3000,

Molecular Probes), mouse anti-22C10 (1:10, Develop-

mental Studies Hybridoma Bank (DSHB), University of

Iowa, IA), rat anti-Elav (7E8A10) (1:10, DSHB), mouse

anti-Repo (1:100, DSHB), mouse anti-Prospero (1:10,

DSHB), mouse anti-Slit (C555.6D) (1:10, DSHB), mouse

anti-Robo (13C9) (1:10, DSHB), rabbit anti-α-tubulin

85E (1:50, Thermo Fisher Scientific), and rabbit anti-

GFP (1:5000, ThermoFisher Scientific). The following

secondary antibodies were used at 1:500: anti-Mouse

555, anti-Rabbit 488, anti-Rat 488, anti-Rat 647, biotinyl-

ated anti-mouse and biotinylated anti-rabbit.

Imaging

A Nikon Eclipse CiL compound light microscope using
differential interference contrast (DIC) optics was used to
take images of Horseradish Peroxidase- (HRP)-stained
embryos at 20X and 40X magnification. A Nikon Eclipse
Ti Confocal Microscope was used to image fluorescence-

stained embryos at 40X and 60X objective lenses. All the

images were analyzed and processed using Nikon Ele-

ments Ar software.

Statistical analysis

A G-test of statistical independence was used to com-

pare abnormality in lch5 neuron migration patterns as

well as the classes of defects and the percentages.

Results

Absence of sli displays improper migration and irregular

morphology of lch5 neurons

It was mentioned previously, but not shown, that the ab-

sence of sli resulted in mis- or non-migration of the lch5

chordotonal organs [19]. Therefore, we wanted to ex-

amine more closely the lch5 neurons in embryos missing

the extracellular ligand sli. Our experiments show that

indeed, while wild-type embryos displayed lch5 neurons

with a teardrop shape and dendrites facing dorsal-

posterior (Fig. 2A, B, black arrows), the absence of sli

(sli2) showed two striking defects: irregular morphology of

the lch5 neurons (Fig. 2C, D, black arrowhead) and

mis-migrating lch5 neurons (Fig. 2C, D, red arrows). We
defined irregular morphology as the lch5 neurons dis-

playing the following traits: neurons lacking a teardrop

shape, neurons not slightly overlapping, and dendrites

pointing in aberrant directions instead of dorsal-

posterior (compare Fig. 2F to Fig. 2E). When quantified,

sli mutants displayed 30% more lch5 neurons with an ir-

regular morphology compared to wild-type (Fig. 2G).

In the wild-type, the lch5 organ migrates ventrally to

its final position in the lateral cluster of peripheral neu-

rons [15, 17, 18]. We found that sli mutants displayed mis-

migrating lch5 neurons that either do not migrate as a

cluster and end up in various positions along the dor-

sral-ventral axis (Fig. 2C, D, red arrows) or completely

fail to migrate (Fig. 2I, white arrow). This failure to mi-

grate can be seen by the distance between the lch5 clus-

ter and the v’ch1 neuron (compare white arrow (lch5

cluster) to white arrowhead (v’ch1 neuron) in Fig. 2H

and Fig. 2I). When quantified, sli mutants displayed a

significantly greater number of lch5 neurons that mis-

migrated or failed to migrate ventrally to their target

location (Fig. 2I). In order to confirm that both these de-

fects were the result of an absence of sli, we also

knocked down sli using a ubiquitous GAL4 driver,

Act5C-GAL4 along with UAS-dcr2 (dicer2) to facilitate

the RNA interference (RNAi). Surprisingly, knocking
down sli displayed a similar, but less severe lch5 neu-

ronal defects than the sli2 null allele (Fig. 2G and J). We

interpret this to mean that even a small amount of Sli

protein present can prevent the lch5 neuronal defects.

Clearly, these data suggest that Sli secreted into the

ECM surrounding these PNS neurons plays an import-

ant role in their migration and final patterning. We next

wanted to know whether two Sli receptors, Robo and

Robo2, play a role in lch5 neuronal morphology and

migration.

Sli is expressed in the epidermis while Robo is expressed

on lch5 neurons

Before examining robo and robo2 mutants for potential lch5

defects, we wanted to confirm the expression patterns of Sli

and Robo. It has been shown that Sli protein gets secreted

from epidermal cells [17, 19]. We wanted to examine more
closely the expression of Sli in relation to the lch5 neurons.

Therefore, we used an epidermal-specific GAL4 driver (69B-

GAL4) to express a secreted GFP (UAS-secGFP) and

compare the expression of the secreted GFP to Sli, and compare

Sli to the lch5 neurons (Fig. 3A–A’). Sli expression (red)

overlaps with the GFP secreted from the epidermal cells (green)

(compare Fig. 3A’ to 3A”) and is not expressed in the

lch5 neurons (blue) (white arrows in Fig. 3A and A”). It

has previously been shown that Robo is expressed at the tips
Fig. 2 (See legend on next page.)
Fig. 2 Absence of sli results in mis-migrating and morphological irregularities in lch5 chordotonal neurons. 

a. WT stage 16 embryo labeled with anti-22C10 to mark all PNS sensory neurons showing the lch5 cluster in a lateral position in the embryo and the five lch5 neurons in an evenly-spaced diagonal row facing the dorsal-posterior edge of the embryo. 

b. Blown-up view of boxed region in A showing three abdominal hemisegments of PNS neurons. Black arrows point to lch5 neurons. 

c. sli mutant (sli2) embryo showing lch5 neurons either mis-migrating or displaying an irregular morphology. 

d. Blown-up view of boxed region in C showing three abdominal hemisegments of PNS neurons. Red arrows show lch5 neurons that have mis-migrated, and black arrowhead shows lch5 neurons with an irregular morphology. 

e. One hemisegment of a WT embryo labeled with anti-22C10 (red) which marks the cell membranes of all PNS neurons and anti-Elav (green) which stains the nuclei of all neurons. White arrow points to the lch5 neurons with normal morphology and the white arrowhead points to the v'ch1 neuron which is normally just ventral to the lch5 cluster. 

f. sli2 embryo showing lch5 neurons with irregular morphology (white arrow). The white arrowhead points to the v'ch1 neuron which is normally just ventral to the lch5 cluster. 

g. Quantification of percent total lch5 clusters displaying irregular morphology in WT, sli2, and a ubiquitous knock-down of sli using RNAi (Act5C-GAL4::UAS-sliRNAi) abbreviated as Act5C::sliRNAi. 

h. WT embryo showing lch5 cluster that has migrated to the correct lateral position. White arrow points to the position of the lch5 neurons and the white arrowhead points to the v'ch1 neuron which is normally just ventral to the lch5 cluster. 

i. sli2 embryo showing lch5 neurons that have not migrated ventrally (white arrow). The white arrowhead points to the v'ch1 neuron. The increased distance between the lch5 neurons and the v'ch1 neurons is visualized in the sli2 compared to the WT in h. 

j. Quantification of percent total lch5 mis-migrating clusters in WT, sli2, and Act5C::sliRNAi. Number of abdominal lch5 clusters scored for each genotype above each bar (n). **p < 0.05 based on a G-test of statistical independence.

Fig. 3 Sli is expressed in the epidermis while Robo is expressed on lch5 dendrites. A-A′. Epidermal 69B-GAL4 was used to drive a secreted GFP (UAS-secGFP) (69B-GAL4::UAS-secGFP) and labeled with anti-Sli (red) (A′), anti-GFP (green) (A″), and anti-Elav (blue) (A‴). White arrows point to lch5 neurons (A and A‴) and white arrowheads point to muscle attachment sites (MASs) expressing Sli (A and A‴). 

B-B‴. WT embryo labeled with anti-Robo (red) (B‴) and anti-Elav (green) (B‴). White arrows point to lch5 neurons and blue arrows point to lch5 dendrites expressing Robo.
of the lch5 dendrites [17]. Here we confirm that Robo (red) is indeed expressed at the tips of the lch5 dendrites (Fig. 3A, B, blue arrows), as well as faintly along the membrane of the lch5 neuronal cell bodies (green) (Fig. 3A, C, white arrows). These results suggest the possibility that Robo acts as the receptor for Sli in the migration and positioning of the lch5 chordotonal neurons.

Absence of robo or robo2 result in mis-migrating and morphological irregularities in lch5 chordotonal neurons

Based on the reciprocal expression patterns of Sli secreted from the epidermal cells and Robo on the lch5 neurons, we wanted to know whether the absence of robo would display any lch5 neuronal defects similar to those observed in the absence of sli. Embryos mutant for robo (robo570) displayed a significant increase in lch5 neurons with an irregular morphology (Fig. 4C, D, G, black arrowheads in 4D). The absence of robo also displayed a slight, but significant, increase in mis-migrating lch5 neurons (Fig. 4H). Interestingly, embryos mutant for robo2 (robo2x135) also displayed a significant increase in lch5 neurons with an irregular morphology (Fig. 4G). Surprisingly, the absence of robo2 displayed almost twice as many mis-migrating lch5 neurons as robo mutants (Fig. 4E, F, H, red arrow in 4F). In this case, the lch5 neurons are too far dorsal as compared to the v’ch1 neuron (white arrowhead in Fig. 4F). These results, as well as previous work, suggests that Robo and Robo2 act together as receptors for Sli to guide and properly position the lch5 chordotonal neurons [19].

Attachment and cap cells follow aberrant lch5 dendrites in absence of sli

Knowing that the lch5 chordotonal neurons are surrounded by several different types of glial cells, we wanted to know how loss of sli affects the position of these glial cells. The ligament cells, cap cells, and attachment cells all express the molecular marker α-tubulin85E [18], shown in a cartoon diagram (Fig. 5A, green cells) compared to the lch5 neurons (Fig. 5A, red cells). Of these specific glial cells, only the ligament cells come into direct contact with the lch5 neurons (Fig. 5A, ventral green cells) [18, 22].

Fig. 4 Absence of robo or robo2 result in mis-migrating and morphological irregularities in lch5 chordotonal neurons. a. WT stage 16 embryo labeled with anti-22C10 showing the lch5 cluster in a lateral position in the embryo and the five lch5 neurons in an evenly-spaced diagonal row facing the dorsal-posterior edge of the embryo. b. Blown-up view of boxed region in A showing two abdominal hemisegments of PNS neurons. Black arrows point to lch5 neurons and white arrowheads point to the v’ch1 neuron, just ventral to the lch5 cluster. c. robo mutant (robo570) embryo showing lch5 neurons displaying an irregular morphology. d. Blown-up view of boxed region in C showing two abdominal hemisegments of PNS neurons. Black arrowheads show lch5 neurons with an irregular morphology and white arrowheads point to the v’ch1 neuron. e. robo2 mutant (robo2x135) embryo showing lch5 neurons not migrating as far ventral. f. Blown-up view of boxed region in E showing two abdominal hemisegments of PNS neurons. Red arrow shows lch5 neurons that are further from the v’ch1 neuron (white arrowhead). g. Quantification of percent total lch5 clusters displaying irregular morphology in WT, robo570, and robo2x135. h. Quantification of percent total mis-migrating lch5 clusters in WT, robo570, and robo2x135. Number of abdominal lch5 clusters scored for each genotype above each bar (n). **p < 0.05 based on a G-test of statistical independence.
Although the ligament cells were harder to visualize with α-tubulin85E, we did observe tubulin filaments (Fig. 5B, white arrowhead) dorsal to the lch5 neurons pointing in the same direction as the dendrites in wild-type embryos (Fig. 5B, white arrow). In sli mutants when the lch5 neurons have mis-migrated or displayed an irregular morphology, tubulin filaments (Fig. 5C, white arrowhead) are seen pointing in the same direction as the irregularly pointed dendrites (Fig. 5C, white arrow). These data suggest that the support cells dorsal to the lch5 neurons, the cap and attachment cells, follow the irregularly pointed dendrites on the lch5 neurons in the absence of sli.

**Scolopale cells follow mis-migrating lch5 neurons in absence of sli**

The scolopale cells are glial cells that are dorsal to the lch5 neurons and in direct contact with the dendrites of the lch5 neurons (Fig. 6A, scolopale cell in red, lch5 neurons in green). We have already seen that the cap and attachment cells (glial cells that are also dorsal to the lch5 neurons) follow aberrantly pointing dendrites of the lch5 neurons in the absence of sli, we asked whether the scolopale cells are also following the mis-migrating lch5 neurons in the absence of sli. To answer this question, we used a marker specific for scolopale cell nuclei, Prospero. In wild-type, the scolopale cells are dorsal to the lch5 neurons (Fig. 6B, white arrow). For this experiment we labeled the lch5 neurons with anti-Elav (a pan-neuronal nuclear marker), and therefore, were not able to visualize the lch5 dendrites. However, in the absence of sli the lch5 neurons are not positioned properly in relation to each other (compare the lch5 neurons in Fig. 6C, white arrowhead with lch5 neurons in 6B, white arrowhead), and the scolopale cells are still dorsal to these neurons (Fig. 6C, white arrow). Additionally, when the lch5 neurons have not migrated fully, the scolopale cells continue to associate with the lch5 dendrites (Fig. 6D, white arrows). These data suggest that the scolopale cells also follow the mis-migrating and irregularly shaped lch5 neurons in the absence of sli.

**Ligament cells continue stretching ventrally in the absence of sli**

The ligament cells are ventral to the lch5 neurons and stretch ventrally from stage 15 (Fig. 7A, red cells) to stage 16 (Fig. 7D, red cells) [18, 22]. We asked whether the ligament cells are coordinated with the lch5 neurons, even when they have mis-migrated or do not orient themselves correctly, as seen in the absence of sli (Fig. 2). At stage 15, wild-type ligament cells can be seen directly ventral to the lch5 neurons (Fig. 7B, white arrow). However, in sli mutants where the lch5 neurons are not in the normal orientation, the ligament cells are either ventral or dorsal to the lch5 neurons (Fig. 7C, white...
arrows). It is not possible to know whether the ligament cells are dorsal or ventral to the lch5 neurons because we are using a nuclear marker for the neurons and, therefore, cannot see where the dendrites of the lch5 neurons are compared to the ligament cells. However, we have never observed an lch5 neuron with its dendrite pointing ventrally. At stage 16, the ligament cells in wild-type start stretching ventrally and have a more elongated shape (Fig. 7E, white arrow). Interestingly, the same stretching of the ligament cells ventrally with a more elongated shape is seen in sli mutants (Fig. 7F, white arrow). These data suggest that the ventral stretching of the ligament cells is independent of mis-migrating lch5 neurons.

Discussion

In this study we show that the extracellular ligand sli is required for the ventral migration and morphology of the lch5 chordotonal neurons in the Drosophila embryonic PNS (Fig. 2). The absence of sli results in lch5 neurons that do not migrate ventrally enough or that have aberrantly pointing dendrites. Interestingly, the absence of the Sli receptors robo and robo2 display similar lch5 neuronal defects to that of sli mutants (Fig. 4), which adds another role to Slit-Robo signaling in neural development. Moreover, we examined the glial support cells that surround the lch5 neurons and make up the lch5 organ. We found that the dorsal glial cells (scolopale, cap, and attachment cells) followed the mis-migrated neurons in the absence of sli (Figs. 5, 6). Conversely, the ligament cells, which are ventral to the lch5 neurons, seem to be independent of the lch5 neurons, as they continue stretching ventrally even when the lch5 neurons mis-migrated (Fig. 7). These results provide further evidence for the possible role for Slit-Robo signaling in lch5 neuronal migration and positioning.

Slit-Robo signaling in the PNS

The Slit-Robo signaling pathway plays a critical role in the development of several organ systems, either as repulsion or attraction [5, 10, 11, 17, 19, 29–32]. In the CNS, Sli is secreted by the midline glia, and the absence of Slit-Robo signaling results in longitudinal axons over-migrating by re-crossing the midline due to a complete lack of repulsive signaling [8–11, 33]. In the embryonic PNS, Sli is expressed in the ektoderm [19] as well as in the visceral mesoderm [17], and secreted into the extracellular environment to interact with various Robo and Netrin receptors to guide neurons and axons [17, 19, 34]. In the thoracic segments of the PNS, the absence of Slit-Robo signaling resulted in thoracic chordotonal neurons migrating ventrally when they normally do not migrate, meaning that Slit-Robo signaling normally inhibits thoracic chordotonal neuron migration [17]. However, in this study we
focused on the abdominal segments, and in the absence of \textit{sli}, the abdominal chordotonal neurons (Ich5) did not migrate ventrally. This result could mean that Sli is acting as a repellant, same as in the thoracic segments. The expression of Robo and Robo2 on the dendrites of the Ich5 neurons [17, this study] could mean that the interaction between the Robo receptors on the Ich5 neurons and Sli in the extracellular space dorsal to the Ich5 neurons, either dorsal or ventral to the Ich5 neurons (white arrows). The absence of either Robo receptor alone displays a defect similar, but less severe, to \textit{sli} mutants. Therefore, a double mutant of \textit{robo} and \textit{robo2} could show a much closer defect to \textit{sli} mutants, as has been mentioned [19]. If Robo and Robo2 were not expressed on the dendrites that are facing dorsal, it might be a possibility that Slit-Robo signaling might be acting as an attraction, as in the somatic muscles [29].

Although a significant number of Ich5 neurons mis-migrate, the more prevalent defect we observed was an irregular morphology, namely aberrantly pointing dendrites in the absence of Slit-Robo signaling (Figs. 2 and 4). Two possibilities arise as an explanation for this defect: (1) if Sli is not present, Robo and Robo2 have no attractive signal to help the Ich5 neurons position themselves properly, or (2) if Sli is not present, Robo and Robo2 have no repulsive signal to stay away from, and as a result, the dendrites are pointing in all directions. Both possibilities are plausible, however, the mis-migration defect leans toward Slit-Robo signaling acting in a repulsive way. Over-expression analysis of \textit{sli}, \textit{robo} and \textit{robo2} would need to be done to figure out the exact mechanism of \textit{sli} action in the migration and rotation of the Ich5 neurons.

**Dorsal glial cells and the Ich5 chordotonal organ**

In the absence of \textit{sli}, the dorsal secondary support cells, scolopale, cap, and attachment cells, seem to follow the mis-migrating Ich5 neurons and aberrantly pointing dendrites (Figs. 5 and 6), which brings up the question of the role of these particular glial cells in the final formation of the Ich5 chordotonal organs. If these dorsal glial cells were independent of the Ich5 neurons and responded to other signals, we might expect these glial cells to migrate and position themselves correctly in \textit{sli} mutants. However, there may be signals coming from the Ich5 dendrites that signal to the scolopale, cap, and attachment cells to stay with or follow the neurons, even if the neurons do not migrate or position themselves properly. It would be interesting to examine these glial cells in other mutants where the Ich5 neurons mis-migrate. A recent report in \textit{C. elegans} describes one of the functions of glial cells associated with sensory neurons is to control the shape of the neuronal endings...
This study brings up the issue that glial cells are not passive, but instructive, especially in the final positioning and functioning of sensory neurons. Therefore, it would also be interesting to observe lch5 neuron migration and positioning in the absence of the glial cells.

Ligament cells and ventral migration of lch5 chordotonal organs

Neurons and glial cells function together in development of the nervous system. We, as well as others, have shown when sli is missing the ligament cells do not have any attachment to the neurons and continue to stretch ventrally on their own (Fig. 7) [17, 18, 22]. This result does not definitively indicate whether the ligament cells are independent of the lch5 neurons, only that they may also be responding to an attractive cue from the ventral ectoderm. Additionally, the absence of sli does not give us the following information: (1) which signal the ligament cells are responding to, (2) whether the signal is attractive or repulsive, and (3) where is the signal coming from. Much more work would need to be done to characterize the ligament cells and what cell surface receptors they express.

Conclusion

In conclusion, our study suggests an added role for Slit-Robo signaling in the development of the nervous system, namely in migration and positioning of the lch5 chordotonal neurons. Although the defects observed in the absence of Slit-Robo signaling are not severe, they are clear, and point to the idea that several different signaling pathways must function together to allow for proper development of any organ system. Clearly, this study opens the door for additional work on the unique properties of the lch5 organ.

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Authors’ contributions

MG and AI designed all the experiments. MG and AI performed all the experiments. MG and AI acquired all the microscopy images. MG and AI conducted all the statistical analyses. MG and AI wrote the paper. All authors read and approved the final version of the manuscript.

Authors’ information

AI is an Assistant Professor at a Primarily Undergraduate Institution, and her research lab is made up solely of undergraduate students. MG was an undergraduate when she performed the experiments for this study.

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