**ccdc80-l1** Is Involved in Axon Pathfinding of Zebrafish Motoneurons

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**Abstract**

Axon pathfinding is a subfield of neural development by which neurons send out axons to reach the correct targets. In particular, motoneurons extend their axons toward skeletal muscles, leading to spontaneous motor activity. In this study, we identified the zebrafish Cccdc80 and Cccdc80-like1 (Cccdc80-l1) proteins *in silico* on the basis of their high aminodic sequence identity with the human CCDC80 (Coiled-Coil Domain Containing 80). We focused on cccdc80-l1 gene that is expressed in nervous and non-nervous tissues, in particular in territories correlated with axonal migration, such as adaxial cells and muscle pioneers. Loss of cccdc80-l1 in zebrafish embryos induced motility issues, although somitogenesis and myogenesis were not impaired. Our results strongly suggest that cccdc80-l1 is involved in axon guidance of primary and secondary motoneurons populations, but not in their proper formation. cccdc80-l1 has a differential role as regards the development of ventral and dorsal motoneurons, and this is consistent with the asymmetric distribution of the transcript. The axonal migration defects observed in cccdc80-l1 loss-of-function embryos are similar to the phenotype of several mutants with altered Hedgehog activity. Indeed, we reported that cccdc80-l1 expression is positively regulated by the Hedgehog pathway in adaxial cells and muscle pioneers. These findings strongly indicate cccdc80-l1 as a down-stream effector of the Hedgehog pathway.

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**Introduction**

The development of a functional vertebrate nervous system requires elaboration of a large number of diverse cell types. At embryonic stages, the nervous system is a complex network of growing axons, whose growth cones navigate in response to guidance cues. Among them, motoneuron axons migrate towards skeletal muscles, and form synaptic contacts [1]. Zebrafish embryos exhibit spontaneous contractions of the musculature ever since 18–19 hours post fertilization (hpf) when removed from their protective chorion [2]. These movements are due to the early-developing primary motoneurons (PMNs), that innervate the myotome with nonoverlapping arbor. In zebrafish, PMNs are present in each somitic hemisegement and are identified by their specific axonal pathway and soma position within the spinal cord: caudal primary motoneurons, middle primary motoneurons and rostral primary motoneurons (CaPs, MiPs and RoPs, respectively) [3,4,5]. PMNs extend their axons out of the spinal cord at about 16–17 hpf, following a common pathway: their growth cones project ventrally along the medioventral surface of the myotome and pause at the horizontal myoseptum, which separates dorsal and ventral myotomes. Here, they specifically interact with muscle pioneers [6,7], a subset of two to six cells for each somite early differentiating into slow muscle fibers [8,9]. CaPs are responsible for pioneering the common pathway before projecting the axons that innervate the ventral myotome [10]. Among PMNs, they show the largest and most extensive branching pattern [5]. MiPs sprout a collateral axon to innervate dorsal myotome, while the first ventral process extending along the common pathway is retracted by 48 hpf [7]. RoPs innervate the middle region of the muscle segment, sprouting laterally after pausing at the myoseptum [6]. Therefore, muscle pioneers represent a choice point from which motoneurons select their specific pathway, although the ablation of this cell population leads to abnormal motor axonal extension without altering the target choice [7]. Secondary motoneurons (SMNs) growth cones extend later from spinal cord, beginning at 22 hpf and following the paths pioneered by the primary axons [11,12].

Axonal pathfinding is dependent on attractive and repulsive stimuli coming from both nervous and non-nervous surrounding tissues [1,13]. For instance, *shh* induction by notochord and floorplate patterns both primary and secondary motoneurons [14]. Indeed, mutants lacking both the notochord and the floorplate (zyg “flh”) [14] or mutants in the Hedgehog pathway, such as *smoothened* (smo), present disorganized, reduced or absent PMNs and axons. [15]. Moreover, in *sonic-yo* mutants (syw) CaPs and MiPs axons run along the neural tube horizontally instead of ventrally and dorsally, while axons of the secondary motoneurons fail to branch and instead cease to extend or grow further ventrally in an abnormal pattern [16].

Also muscular tissues can pattern axonal migration: muscular adaxial cells are able to rescue motor axon defects in *dicycledon*...
mutants, showing that this myotomal pattern plays a pivotal role in axonal migration [17]. Furthermore, molecules expressed in the somites such as the semaphorin proteins Zib and Sem3A1 are involved in repelling axonal migration [10,16]. On the contrary, netrin-1a is expressed by adaxial cells and muscle pioneers besides ventral spinal cord, and seems to guide axonal growth through a chemotactic function [19]. The manipulation of the proper expression of these molecules, both knocking-down and inducing ectopic expression, induces axons to follow aberrant pathways, branch excessively or stall [17,18,19].

The Coiled-Coil-Domain Containing 80 (Ccd80) gene, also named DRO1 in human, Down-Regulated by Oneogene (Down-Regulated by Oneogene), URB in mouse (Up-Regulated in BRS-3-deficient mice), CL2 in rat (Clone 2), and squarin in chicken, has been recently suggested to be involved in different functions among vertebrates. Ccd80 was first isolated in mice, where it is up-regulated in adipose tissue of obese BRS-3-deficient animals [20]. Moreover, Ccd80 is highly expressed in mice white adipose tissue and its silencing inhibits adipocytes differentiation [21], suggesting a role in the regulation of body weight and energy metabolism. Ccd80 is also expressed in mouse developing cartilage, suggesting a role during skeletogenesis [22].

Human CCDC80 is almost ubiquitously expressed, with the highest levels in heart and skeletal muscles [23,24]. Furthermore, human CCDC80 can be considered a potential tumor suppressor gene [25]. In fact, it is strikingly down-regulated in thyroid cancer cell lines and tissues, as well as in colon and pancreatic cancer cell lines and in most colorectal cancer specimens [26], while its ectopic expression in these cell lines results in substantial inhibition of growth properties.

The CCDC80 protein is highly conserved among vertebrates, and contains multiple signals of cellular compartmentalization and post-translational modifications. In particular, it has a N-terminus leader peptide for extracellular export and many nuclear localization signals [25]. In different studies, the CCDC80 protein has been identified in a N-glycosylated form and was suggested to be secreted. Rat, mouse and human CCDC80 show three P-DUDEs (Procyotyes) (DRO1-URB-DRS-Squarin-SRPUL) which in human are correlated with a tumor suppressor role [27].

In silico analysis using human CCDC80 sequence as a bait, led to the identification of three zebrafish homologs of CCDC80. Two homologs, that we named Ccd80 and Ccd80-like1 (Ccd80-l1), have high levels of aminocacid identity with the human CCDC80. We performed the molecular cloning of Ccd80 and ccd80-l1 in zebrafish and analyzed their expression pattern during embryonic development. During somitogenesis ccd80 is expressed in the notochord (manuscript in preparation), while ccd80-l1 is expressed in muscle pioneers and adaxial cells. Both these regions are responsible for axon guidance, therefore we decided to investigate the role of ccd80 and ccd80-l1 in this process. While loss-of-cdd80 function did not impair motorneurons development, we demonstrated the ccd80-l1 involvement in the proper axonal pathfinding, especially in ventral axons guidance. Indeed, ccd80-l1 knocked-down embryos exhibited motility issues although analysis on body musculature showed that somitogenesis and myogenesis occurred properly. Furthermore, the analysis of ccd80-l1 upstream regulation revealed that the Hedgehog pathway modulates its expression in territories involved in axonal guidance.

Materials and Methods
Zebrasfish lines and maintenance
Current Italian national rules: no approval needs to be given for research on zebrafish embryos. Zebrafish were raised and maintained according to established techniques [28], approved by the veterinarian (OVSAC) and the animal use committee (IACUC) at the University of Oregon, in agreement with local and national sanitary regulations. Embryos were collected by natural spawning, staged according to Kimmel [29], and raised at 28°C in fish water (Instant Ocean, 0.1% methylene blue) in Petri dishes [30].

Sequence analysis
Zebrafish chromosome 6 region hosting the ccd80-l1 gene was identified through in silico search of the ENSEMBL zebrafish assembly version 9 (Zv9) using human CCDC80 aminoacidic sequence as a bait. The alignments between aminoacidic sequences were performed with the software program StrecherP. Analysis on synteny was performed with the program Genomicus version 57.01.

RT-PCR
Total RNA from 11 samples (an average of 30 embryos per sample) corresponding to 9 different developmental stage embryos (2–4 cells, 64–1000 cells, 30% epiboly, 60%–70% epiboly, somitogenesis, 24 hpf and 72 hpf) and 2 adult organs (ovary and muscle) was extracted with the TOTALLY RNA isolation kit (Ambion), treated with RQ1 RNase-Free DNase (Promega) and oligo (dT)-reverse transcribed using Super-Script II RT (Invitrogen), according to manufacturers’ instructions. The following primers were used for PCR reactions: ccd80-l1_forward 5'- ACCACAATTTGAGCAAACAACA -3' and ccd80-l1_reverse 5'- GGTTATTAGCCTCCTCCCTTGG -3'. PCR products were loaded and resolved onto 2% agarose gels.

In situ hybridization and immunohistochemistry
Whole-mount in situ hybridization (ISH), was carried out as described [31] on embryos fixed for 2 hours in 4% paraformaldehyde in PBS, then rinsed with PBS-Tween (PBT), dehydrated in 100% methanol and stored at −20°C until processed [32]. Antisense riboprobes were previously in vitro labelled with modified nucleotides (digoxigenin, Roche). nycd and nycg probes were prepared as described by Schnapp and colleagues [33]. snahe1 probe has been kindly provided by Ingham laboratory. The following primers were used for PCR reactions to clone the probes: ccd80-l1 sense 5'- ACCACAATTTGAGCAAACAACA -3' and ccd80-l1 antisense 5'- GGTTATTAGCCTCCTCCCTTGG -3'. For immunohistochemistry, embryos were fixed in 4% paraformaldehyde overnight at 4°C or 2 hours at RT, washed several times in PBT and blocked in 5% BSA in PBT for 1 hour at room temperature. Primary antibody incubation was done overnight at 4°C, followed by several washes in PBT and incubation of secondary antibody for 1 hour at room temperature. Primary antibodies are MF20 (mouse anti-sarcomeric) and 4D9 (mouse anti-engrailed/invected) purchased from Developmental Studies Hybridoma Bank, znp1 (mouse anti-syphilis) and zn-5 (mouse anti-alkaline) purchased from Zebrafish International Resource Center (ZIRC). Secondary antibody is EnVision+ System- HRP Labelled Polymer anti-mouse (Dako). Images of embryos and sections were acquired using a microscope equipped with a digital camera with LAS Leica imaging software (Leica, Wetzlar, Germany). Images were processed using the Adobe Photoshop software. For histological sections, stained embryos were re-fixed in 4% paraformaldehyde, dehydrated and stored in methanol, wax embedded, and sectioned (3–8 μm).

Injections
Injections were carried out on 1- to 2-cell-stage embryos (with Eppendorf Femtotjet Micromanipulator 5171); the dye tracer rhodamine dextran was co-injected as a control. To repress ccd80-l1 mRNA translation we designed an ATG-targeting morpholino.
The table shows the scores obtained after alignments between the aminoacidic sequences of zebrafish and human CCDC80 homologs. Alignments were performed with Stretcher-P tool.

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Table 1. Percentages of identity and similarity among human and zebrafish Ccd80 homologs.

|                      | Human CCDC80 | Zebrafish Ccd80 | Zebrafish Ccd80-like1 | Zebrafish Ccd80-like2 |
|----------------------|--------------|-----------------|----------------------|----------------------|
|                      | Identity    | Similarity     | Identity    | Similarity     | Identity    | Similarity     | Identity    | Similarity     |
| Zebrafish Ccd80      | 51.6%       | 65.2%          | /           | /             | 51.4%       | 64.4%          | 30.4%       | 47.4%          |
| Zebrafish Ccd80-like1| 44.4%       | 59.3%          | 51.4%       | 64.4%         | /           | /             | 29.1%       | 46.9%          |
| Zebrafish Ccd80-like2| 27%         | 44.9%          | 30.4%       | 47.4%         | 29.1%       | 46.9%          | /           | /              |

cdc80-ll-MO: 5’- TTTGACCTGTAGATTTTGTACCA-3’

and a splice-site morpholino, cdc80-ll-spliace- 5’- TGATACATAA

CATACTGAGGCCTGTG-3’ (Gene Tools, LLC). As a negative

control we injected a standard control morpholino oligonucleotide

(ctrl-MO). Morpholinos were injected in 1 x Danau buffer (pH 7.6)

as suggested by Nasevicius and Ekker [34]. For the in vivo test of the

efficiency of cdc80-ll-MO, 425 pg/embryo of the pCS2+-cdc80-ll-

GFP sensor plasmid has been injected alone or co-injected with

12 ng/embryo of cdc80-ll-MO. The presence/absence of the GFP

signal has been monitored under a fluorescent microscope starting

12 ng/embryo of GFP sensor plasmid have been injected alone or co-injected with

as suggested by Nasevicius and Ekker [34]. For the

we named

chromosome 6 (nucleotide position: 16,322,724-16,342,517) that

(ccdc80-like2

(nucleotide position: 18,662,129-18,669,986), that we named

incubations in 0.05% ethanol in embryo medium.

PFA at 15 somites stage. Cyclopamine was dissolved in embryo

SIGMA-ALDRICH) from 50% epiboly stage up to fixation in

motility

To determine the functional role of cdc80-ll during zebrafish
development, we specifically knocked it down by means of the

injection of an antisense oligonucleotide morpholino (cdc80-ll-MO, Gene Tools) designed against the start site of the transcript. In all the experiments, cdc80-ll-MO-injected embryos (morphants) were compared to embryos at the same developmental stage, injected with the same amount of a control MO (ctrl-MO). For the in vivo test of the efficiency of cdc80-ll-MO, 425 pg/embryo of the pCS2+-cdc80-ll-GFP sensor plasmid has been injected alone or with 12 ng/embryo of cdc80-ll-MO. The presence/absence of the GFP signal has been monitored under a fluorescent microscope starting from somitogenesis up to 48 hpf (Fig. S2). Moreover, analysis of chromosomal organization of the three cdc80 zebrafish homologs across vertebrates revealed that only cdc80 is syntenic with other vertebrates (Fig. 1).

cdc80-ll is expressed in muscle pioneers and adaxial cells of the zebrafish embryo

Characterization of cdc80-ll expression, using RT-PCR, revealed that cdc80-ll transcript is present from the first stages of development up to 72 hpf, thus including maternal and zygotic transcription (Fig. 2A). cdc80-ll is also expressed in the ovary and muscle of the adult zebrafish (Fig. 2A). During somitogenesis, the hybridization signal is restricted to the horizontal myoseptum (Fig. 2B–D). From this stage, cdc80-ll expression is observed also in the cranial ganglia and dorsal dermis (Fig. 2B, 2C, 2E, 2F, 2K–L and data not shown). At 24 hpf, cdc80-ll is detectable in a specific sub-population of migrating adaxial cells, that moves along the lateral axis towards the external somite [35] (Fig. 2E–G). Moreover, cdc80-ll is expressed in muscle pioneers, as shown by the co-localization between cdc80-ll and engulfed cell (36,37) (Fig. 2H). cdc80-ll expression in adaxial cells persisted at 36 hpf and 48 hpf (Fig. 2I, 2K). At the same stages cdc80-ll is also expressed in the caudal veil plexus region (Fig. 2I, 2J, 2L).

cdc80-ll knocked-down embryos displayed impaired motility
loss-of-function was not so striking, as demonstrated by the low percentage of embryos with GFP decreasement and by the high amount of morpholino we had to inject to obtain a phenotype (8 and 12 ng/embryo of ccdc80-l1-MO). Therefore, we designed a second morpholino against the splice site (ccdc80-l1-splice-MO) to confirm the specificity of ccdc80-l1-loss-of-function. Embryos injected with this second morpholino, still exhibited motility issues as ccdc80-l1-MO-injected embryos did (data not shown). In particular, all the knocked-down embryos showed no severe body plan alteration when observed in vivo, indicating the proper

Figure 1. Analysis of chromosomal organization of the three ccdc80 zebrafish homologs across vertebrates. Each ccdc80 gene is shown as a reference locus. Genes annotated as paralogs (no surrounding line) or orthologs (with a surrounding line) by the Ensembl database share the same color, blue lines beneath individual tracks indicate that orientations of gene blocks and are inverted with respect to their genomic annotation. For zebrafish ccdc80 (chr. 9), ccdc80-l1 (chr. 6) and ccdc80-l2 (chr. 21), only ccdc80 shows notable synteny with other vertebrates. The figure was derived from the output of the Genomicus website (version 57.01).

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Figure 2. Expression of ccdc80-l1 analyzed by RT-PCR and WISH. (A) RT-PCR performed on different embryonic stages and adult tissues; the expression of ccdc80-l1 and β-actin are shown. Lanes are: ladder (lane 1), ovary (lane 2), 2-4 cells stage (lane 3), 64-1000 cells stage (lane 4), 30% epiboly (lane 5), 60-70% epiboly (lane 6), somitogenesis (lane 7), 24 hpf (lane 8), 30 hpf (lane 9), 48 hpf (lane 10), 72 hpf (lane 11), adult muscle (lane 12) and negative control (lane 13) in the absence of cDNA. (B–J) WISH performed on zebrafish embryos at several stage of development. (B) During somitogenesis ccdc80-l1 was expressed by cranial ganglia (cg), dorsal dermis (asterisk), adaxial cells and muscle pioneers at the level of the horizontal myoseptum (arrow). (C) ccdc80-l1 expression in a transverse section of the trunk of an embryo at 12 somites stage (arrows). (D–I) At 24 hpf, the hybridization signal was detectable in cranial ganglia (cg), dorsal dermis (asterisk), adaxial cells (arrow) and ventral somites (arrowhead). (F) Higher magnification of the tail at 24 hpf. (G) Transversal section of an embryo at 24 hpf. (H) Transversal section showing that at 24 hpf ccdc80-l1 hybridization signal co-localized with the nuclear labeling of 4D9 antibody, corresponding to the engrailed-positive muscle pioneers population (open arrowhead). (I, J) At 36 hpf, the signal of ccdc80-l1 probe was detected in cranial ganglia (cg), migrated adaxial cells (arrow), dorsal dermis (asterisk) and caudal vein plexus region (cvp). (K, L) At 48 hpf, ccdc80-l1 was detected in dorsal dermis (asterisk), external adaxial cells (arrows in K) and caudal vein plexus region (cvp in L). (B, E, F, I) Lateral views; dorsal is up, anterior is left; (C) dorsal view, anterior is left; (D, G, H, J–L) transversal sections, dorsal is up.

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progression of early developmental processes such as gastrulation and segmentation [38,39,40,41]. Moreover, we observed that morphants displayed physiological body contractions upon dechorionation at 24 hpf [42]. However, at 48 hpf, almost 80% of the morphant embryos (N = 37) presented abnormal escaping behavior after tactile stimulation, often resulting in body contractions on the spot or circling behavior (Video S2) rather than a fast escape in the opposite direction to the stimulus (Video S1). The same phenotype was observed also at 5 days post fertilization (5 dpf, data not shown). These results indicated that \textit{ccdc80-l1} loss-of-function affects the swimming behavior of zebrafish embryos and larvae. We were also able to rescue the \textit{ccdc80-l1}-loss-of-function phenotype by means of the injection of the homolog \textit{ccdc80}-full-length transcript. In fact, despite we demonstrated that \textit{ccdc80}-loss-of-function did not affect axonal pathfinding (data not shown), the high degree of conservation between the two homologs allowed rescue of motility (only 42% of rescued embryos presented motility issues in comparison to the nearly 80% of \textit{ccdc80-l1}-MO injected embryos, N = 63).

\textit{ccdc80-l1} loss of function does not affect somitogenesis nor muscle pioneers and adaxial cells formation

To assess whether the phenotype displayed by morphants was due to the impairment of musculature, we examined somitogenesis and myogenesis markers. The expression of \textit{myod} and \textit{myog} [39,43] was not altered in \textit{ccdc80-l1}-MO-injected embryos (Fig. 3A–D). Moreover, the expression of \textit{smyhc1}, a marker of slow-twitch fibers [44], was unaffected as well, notwithstanding the strong expression of \textit{ccdc80-l1} in adaxial cells and muscle pioneers, from which slow fibers develop [9] (Fig. S3A, S3B). In addition, at 24 hpf, myofibers were correctly organized and distributed as shown by the immunohistochemistry with anti-sarcomeric MF20 antibody [45] (Fig. 3E, 3F). Also muscle pioneers, labeled with 4D9 anti-engrailed antibody [36,37] were correctly formed in \textit{ccdc80-l1} morphants (Fig. 3G–H). These results led us to exclude that defects of adaxial cells, muscle pioneers or body musculature formation were responsible for motility issues observed in \textit{ccdc80-l1} knocked-down embryos.
Proper pathfinding and branching of axons are affected in ccdc80-l1-MO-injected embryos

Segmentally repeated motoneurons connect nervous system to somites, as their growth cones exit the spinal cord during embryogenesis and migrate to their appropriate muscle targets, allowing movement [3,5]. Due to motility impairment of ccdc80-l1 morphants, we investigated the morphology of motoneurons performing immunohistochemistry with zn1 (znp1) antibody [46]. For all embryos, we analyzed the trunk region overhanging the yolk extension; defects in at least three motoneurons were enough to consider the embryo as affected. At 48 hpf, in the 84% (N = 33) of morphants injected with 12 ng/embryo of morpholino, axonal pathfinding resulted impaired. 60% of morphants displayed an overall disorganization of both dorsal and ventral motoneurons, that resulted mis-orientated and over-branched (Fig. 4A, 4B). In the 9% of embryos, these defects were observed together with an opposite phenomenon, axonal stalling. In the 12% of morphants only ventral axons resulted mis-orientated and over-branched, whereas in the 3% only the dorsal ones were affected. This phenotype was dose-dependent: when 8 ng/embryo of morpholino were used, a lower percentage of embryos resulted affected (64%, N = 35). Interestingly, at this concentration, only 27% of the knocked-down embryos displayed both ventral and dorsal defective axons, whereas in the 37% of morphants the same defects were detectable in the ventral motoneurons solely (Fig. 4C).

Dorsal axons alone were never affected (Table 2 and Fig. S4). Thus, a striking reduction of Ccdc80-l1 protein amount led to the affection of both ventral and dorsal motoneurons, whereas a lower dose of morpholino is sufficient for ventral axons migration impairment. In order to discriminate whereas loss-of-

Table 2. The phenotype of ccdc80-l1-MO-injected embryos is dose-dependent.

| Dose/type of morpholino | Total percentage of affected embryos (N) | Alteration of both ventral and dorsal axons | Only defective ventral axons | Only defective dorsal axons |
|-------------------------|----------------------------------------|--------------------------------------------|----------------------------|-----------------------------|
| ctrl-MO 12 ng           | 12% (N = 25)                           | 12%                                        | 0%                         | 0%                          |
| ccdc80-l1-MO 12 ng      | 84% (N = 33)                           | 69%                                        | 12%                        | 3%                          |
| ccdc80-l1-MO 8 ng       | 64% (N = 35)                           | 27%                                        | 37%                        | 0%                          |

The percentage of embryos displaying axonal defects decreased from 84% to 64% when a lower dose of morpholino was used. Both ventral and dorsal axonal pathfinding resulted impaired in the 69% of affected embryos when 12 ng/embryo of morpholino were used. After the injection of the lower dose of ccdc80-l1-MO (8 ng/embryo), 27% of affected embryos showed alteration of both ventral and dorsal axons, whereas the 37% displayed only ventral defective axons and dorsal axons alone were never affected.

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Figure 4. Analysis of motoneurons morphology by means of znp1- and zn-5-immunohistochemistry. (A, B) At 48 hpf, using 12 ng/embryo of morpholino, both ventral (arrows) and dorsal axons (arrowheads) were mis-orientated and over-branched in morphants (B) in comparison to control embryos (A). (C) Statistical analysis showing the percentages of the different phenotypes (affected ventral axons, dorsal axons or both) occurring in control embryos and in morphants, when different doses of ccdc80-l1-MO were injected (12 ng/embryo and 8 ng/embryo). Using a lower dose of morpholino (8 ng/embryo), we observed that in a significant percentage of embryos only ventral axons were defective. (D–G) Immunohistochemistry performed at 26 hpf (D, E) and 30 hpf (F, G) confirmed that loss-of-ccdc80-l1-function affects both CaPs (arrows) and MiPs (arrowheads) axonal migration. (H, I) The same analysis performed at 48 hpf using zn-5 antibody revealed that also SMNs axonal migration is impaired in morphants (arrows in I) in comparison to control embryos (H). (A, B; D–I) Lateral flat-mount preparation was applied for a better visualization of the motoneurons. Lateral views of the trunk region overhanging the yolk extension, dorsal is up and anterior is left.
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function impaired PMNs or SMNs, we analyzed the phenotype of morphants also at 26 hpf and 30 hpf, by the time SMNs have just begun extending axons [47], so most of znp1 labeling correspond to PMNs. At this stages, we replicated the same phenotype observed at 48 hpf (Fig. 4D-G and Table 3). Furthermore, by performing an immunohistochemistry at 48 hpf using the specific antibody for SMNs (zn-5, anti-alcama) [47], we observed that also SMNs axons seems to be affected after loss-of-ccdc80-l1-function (Fig. 4H, 4I). Thus, the analysis of the motoneuronal patterning in morphant embryos revealed the lack of proper guidance toward muscle targets, suggesting ccd80-l1 involvement in axonal pathfinding of both PMNs and SMNs.

ccd80-l1 expression is positively regulated by the Hedgehog pathway

Both muscle and motoneurons induction is finely regulated by levels and range of shh expression [9,48,49]. Due to ccd80-l1 expression in adaxial cells and muscle pioneers, we decided to investigate the existence of a ccd80-l1 up-stream regulation Hedgehog-mediated. We modulated shh activity by exposing embryos to cyclopamine, that inhibits the Hedgehog transducer Smoothened (smo) [9,50]. To avoid the complete loss of the territories in which ccd80-l1 is expressed, we chose a concentration of cyclopamine (5 μM) by which muscle pioneers and adaxial cells-derived slow fibers are unaffected, as already described [9] and as we demonstrated by the proper expression of their markers engrailed, myod and myo5c1 respectively. (Fig. S5). A striking down-regulation of ccd80-l1 expression was observed in 72% of the treated embryos in comparison to controls (N = 32) (Fig. 5A, 5B).

Interestingly, this down-regulation was detectable only at the level of myoseptum and somites, whereas the cephalic territories in which ccd80-l1 is expressed were not involved. A similar down-regulation was observed in syu mutants, carriers of a deletion in the gene sonic-you encoding for shh [Fig. 5D–F] [16]. ccd80-l1 signal in adaxial cells was extremely weak or absent in the 35% of mutants, and slightly down-regulated in the 40% of observed embryos (N = 20). Moreover, the overexpression of shh by means of the injection of the full-length transcript (300 pg/embryo), led to the opposite phenotype with an increasing of ccd80-l1 expression in the somites of the 71% of the injected embryos (N = 31) (Fig. 5C). On the contrary, ccd80-l1 loss-of-function did not affect shh expression (Fig. S6). Therefore, these findings suggest that ccd80-l1 is a down-stream target of the Hedgehog pathway.

Discussion

The genetic program underlying axon guidance is not completely defined. Adaxial cells and muscle pioneers are both involved in axonal outgrowth and pathfinding [7,17], even if little is known about the specific proteins and molecular mechanisms acting in this process. ccd80-l1, the novel gene we recently identified in zebrafish, is expressed during embryonic development in muscle pioneers and adaxial cells. Ccd80-l1 was identified, together with its homolog Ccdd80, on the basis of its high aminoacid identity with human CCDC80. However, zebrafish ccd80 and ccd80-l1 do not share the same expression pattern and seems to play different roles. In fact, only ccd80-l1-MO-injected embryos showed a complete loss of expression.

Table 3. Loss-of-ccdc80-l1-function impairs PMNs axonal migration.

| Developmental stage/dose of morpholino | Total percentage of affected embryos (N) | Alteration of both Caps and Mips | Only Caps affected | Only Mips affected |
|----------------------------------------|------------------------------------------|---------------------------------|-------------------|-------------------|
| 26 hpf/ccd80-l1-MO 12 ng               | 54.5% (N = 33)                           | 40%                             | 11%               | 3.5%              |
| 30 hpf/ccd80-l1-MO 12 ng               | 62% (N = 35)                            | 33%                             | 24.5%             | 4.5%              |

Embryos injected with 12 ng/embryo of ccd80-l1-MO were observed also at 26 hpf and 30 hpf. At these stages, affected embryos were 54.5% and 62%, respectively. The percentages of the different phenotypes are listed.

Figure 5. ccd80-l1 is positively regulated by shh. (A–C) ccd80-l1 expression in somites and myoseptum resulted strongly inhibited in embryos treated with 5 μM cyclopamine (asterisks in B), in comparison to control embryos at the same developmental stage (A). By converse, over-expression of shh led to an up-regulation of ccd80-l1 in muscular territories (C). Expression in cranial ganglia (cg) was never perturbed. (D–F) ccd80-l1 resulted slightly down-regulated in the muscles of heterozygous syu+/− mutants (E) in comparison to wild type siblings (D). A strikingly down-regulation was observed in homozygous syu−/− mutants (F). (A–C) Dorsal flat-mount preparations, anterior is up. (D–F) Lateral views of the tails, anterior is left. doi:10.1371/journal.pone.0031851.g005
embryos displayed an abnormal escaping behavior after tactile stimulation at 48 hpf. Both musculature and nervous system are responsible for embryonic motility and touch response and are the basis of spontaneous motor output that occurs in the developing zebrafish embryo ever since 10 hpf [2]. Nevertheless, musculature defects were unlikely the basis of the observed phenotype. Indeed, there was no difference between the expression pattern of myogenic markers in morphants and control embryos. Moreover, muscle fibers resulted correctly formed and distributed by the end of somitogenesis. The territories in which ccd80-l1 is expressed were unaffected as well: in fact, adaxial cells and muscle pioneers showed no defects. These findings revealed that ccd80-l1 function is not necessary for the specification and further differentiation of myogenic cell populations, suggesting that the motility issues displayed by morphants at 48 hpf could be due to an impairment of neuronal development.

The analysis of motoneuronal development in morphant embryos revealed that ccd80-l1 plays a role in motoneurons axonal pathfinding. In fact, ccd80-l1 loss-of-function did not prevent the formation of PMNs and axon projection, but led to an overall disorganization of PMNs. CaPs and MiPs resulted over-branched in a high percentage of embryos, whereas a smaller fraction of morphants displayed also the simultaneous presence of its opposite phenomenon, axonal stalling. Axonal over-branching and stalling were detected in the CaPs solely in a significant percentage of embryos, especially when a lower dose of morpholino was used. Moreover, the impairment of axonal migration was more severe in CaPs then in MiPs, even when both PMNs were affected simultaneously. These data suggest that ccd80-l1 may have a differential role as regards the development of CaPs and MiPs. This is consistent with the asymmetric distribution of ccd80-l1 transcript in the somites: indeed, the ccd80-l1 transcript is present in the ventral portion of somites, innervated by CaPs, and not in their dorsal portion, innervated by MiPs. The same motility issues displayed at 48 hpf were observed also at 5 dpf, when secondary motoneurons are already formed. Therefore, ccd80-l1 plays a role also in guidance of SMNs, as shown by the mis-expression of their marker zn-5. This is not surprising, as the growth cones of SMNs require the axons of PMNs for proper pathfinding [6]. We concluded that the ccd80-l1 loss-of-function prevents the proper development of the peripheral nervous system, that lacks a proper guidance toward muscle target: axons do not follow a single direction-pathway but stall or extend towards any direction, leading to an over-branched and non-functional nervous network. Hence, embryos are able to move and to respond to tactile stimuli, but the coordination of muscle contractions is impaired, and motor behavior is affected.

Axon outgrowth is influenced by many factors, for instance different molecules (netrins, semaphorins, slits) with chemotropic functions (reviewed in [51]) and components of the extracellular matrix (ECM) [52,53]. In fact, the growth cone shares many features with the motile structures of migrating cells, including actin polymerization at the leading edge, dynamic interactions between cell-surface adhesion receptors and components of the extracellular matrix (ECM), and generation of traction forces in the cytoskeleton applied to ECM through adhesion sites [54,55]. In ccd80-l1 knocked-down embryos, the growth cones are still able to exit the spinal cord and reach the muscle pioneers along the common pathway. Moreover, axonal extensions developed without altering the target choice: in fact, CaPs and MiPs still project their axons ventrally and dorsally, respectively. These data are consistent with the proper development of muscle pioneers, which provide a choice point for motor growth cones. However, further defects occur during axon pathfinding. It has been recently reported that DRO1/CCDC30 is a Golgi-associated-protein [56]. Moreover, the in silico prediction of the Ccdc80 protein structure (String 9.0) suggests its interaction with fibronectin, a component of the ECM. If this is the case also for its homolog ccd80-l1, its loss-of-function might interfere with the proper axon migration by influencing the secretion of guidance molecules or by altering interactions with ECM proteins such as fibronectin. Further analysis on the predicted Ccdc80-l1 protein sequence and its interaction with other proteins will be necessary to understand the molecular process underlying ccd80-l1 functioning. Moreover, investigation on possible targets is still needed. For instance, it is to explore the possibility of an interaction with the semaphorin and netrin families, both involved in attracting and/or repelling growth cones from a variety of organisms [13,19]. Nevertheless, our results provide further insights into motoneurons development, a complex mechanism that requires the action of several different molecules. Moreover, we suggest that ccd80-l1 may act as a down-stream effector of shh. The Hedgehog family consists of secreted morphogens fundamental for both axon guidance and formation of adaxial cells and muscle pioneers [9,57]. The Hedgehog signaling is known to play a pivotal role in the specification of both primary and secondary motoneurons [14,49]. Indeed, mutants for different molecules involved in this pathway displayed axonal defects, including random axonal migration or stalling [15,16]. PMNs target choice was never impaired after ccd80-l1 loss-of-function, still axonal migration resulted aberrant. Furthermore, ccd80-l1 expression resulted strikingly down-regulated after exposure to 5 μM of cyclopamine and up-regulated after over-expression of shh. This modulation was observed only in muscles and not in other territories in which s-ccd80 is expressed (cranial ganglia and dorsal dermis). These findings strongly suggest the existence of a specific regulation Hedgehog-mediated of ccd80-l1, as regards its function in motoneuronal development. Moreover, these findings may shed light on the involvement of the Hedgehog pathway in this process.

Supporting Information

Figure S1 Alignment among human CCDC80 and the three zebrafish homologs. * = identical aminoacids; = conservative substitution; . = non-conservative substitution. (TIF)

Figure S2 ccd80-l1 morpholino is capable to inhibit the expression of the fluorescent protein GFP. This assay was performed in order to verify the in vivo efficiency of ccd80-l1-MO. (A, B) In the 70% of embryos injected with the ccd80-l1-GFP sensor plasmid, the presence of fluorescent GFP signal was detected (N = 20). (C, D) When the plasmid was injected together with the morpholino, the transcription of GFP protein was inhibited and the percentage of fluorescent embryos decreased to 51% (N = 93). In A and C embryos are visualized under normal light, in B and D under fluorescent light. (TIF)

Figure S3 The expression pattern of the slow-myosin marker smyhc1 is unaffected in ccd80-l1 knocked-down embryos. (A, B) Loss-of-ccd80-l1-function did not perturb the expression of smyhc1, as morphant embryos (B) are indistinguishable from control embryos (A). Lateral views of the tails, dorsal is up, anterior is left. (TIF)

Figure S4 Statistical analysis of three distinct defects observed after loss-of-ccd80-l1-function. (A–C) The graphics show the occurrence of three axonal migration defects...
in control embryos and morphants when two doses of ccdc80-l1-MO are used: both dorsal and ventral defective axons (A), only ventral defective axons (B) and only dorsal defective axons (C). The last phenotype was not statistically significant. *** p<0.001 vs ctrl-MO. * p<0.05 vs ctrl-MO. (TIF)

Figure S5 Muscle pioneers and adaxial cells are present after 5 μM cyclopamine treatment. (A, B) Labeling with 4D9 antibody (anti-engrailed) showed that muscle pioneers are not missing after pharmacological inhibition of the Hedgehog pathway (arrows). (C–F) Also adaxial cells are still present, as shown by the expression of the markers myod (C, D) and smyhc1 (E, F). (A, B) Lateral views, dorsal is up. (C–F) Dorsal views, anterior is left. (TIF)

Figure S6 shh expression is not perturbed by loss-of- ccdc80-l1-function. (A, B) shh resulted correctly expressed both in control embryos (A) and in morphants (B). (A, B) Dorsal views, anterior is left. (TIF)

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Video S1 Control embryos displayed standard motor behavior. After tactile stimulation, control embryos fast escaped in the opposite direction of the stimulus. (AVI)

Video S2 The motility of morphant embryos is impaired. When morphants were stimulated, abormal escape was observed, also resulting in circling behavior. (AVI)

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Author Contributions

Conceived and designed the experiments: CB AP AF. Performed the experiments: CB AP AF. Analyzed the data: CB AP. Contributed reagents/materials/analysis tools: FS FC. Wrote the paper: CB AP. Contributed to experimental design and provided helpful suggestions: FS IDN. Supervised the research project: FS AP. Supervised paper drafting: CB AP FS.
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