

# Somatic CRISPR–Cas9-induced mutations reveal roles of embryonically essential dynein chains in *Caenorhabditis elegans* cilia

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**C**ilium formation and maintenance require intraflagellar transport (IFT). Although much is known about kinesin-2-driven anterograde IFT, the composition and regulation of retrograde IFT-specific dynein remain elusive. Components of cytoplasmic dynein may participate in IFT; however, their essential roles in cell division preclude functional studies in postmitotic cilia. Here, we report that inducible expression of the clustered regularly interspaced short palindromic repeats (CRISPR)–Cas9 system in *Caenorhabditis elegans* generated conditional mutations in IFT motors and particles, recapitulating ciliary defects in their null mutants. Using

this method to bypass the embryonic requirement, we show the following: the dynein intermediate chain, light chain LC8, and lissencephaly-1 regulate retrograde IFT; the dynein light intermediate chain functions in dendrites and indirectly contributes to ciliogenesis; and the Tctex and Roadblock light chains are dispensable for cilium assembly. Furthermore, we demonstrate that these components undergo biphasic IFT with distinct transport frequencies and turnaround behaviors. Together, our results suggest that IFT-dynein and cytoplasmic dynein have unique compositions but also share components and regulatory mechanisms.

## Introduction

Cilia are microtubule-based structures that play diverse roles in motility, sensory perception, and signaling (Rosenbaum and Witman, 2002; Scholey, 2003; Eggenschwiler and Anderson, 2007; Ishikawa and Marshall, 2011). The assembly and maintenance of cilia require bidirectional intraflagellar transport (IFT), which ferries ciliary precursors bound to the IFT particle protein complex from the base of the ciliary axoneme to the tip (Rosenbaum and Witman, 2002; Scholey, 2003, 2013). Anterograde IFT is powered either by a single heterotrimeric kinesin-2 motor, which was first proposed in *Chlamydomonas reinhardtii*, or by the concerted action of a heterotrimeric kinesin-2 and a homodimeric OSM-3-kinesin in kinesin-2 family in *Caenorhabditis elegans* neuronal cilia (Rosenbaum and Witman, 2002; Scholey, 2003, 2013). Retrograde IFT is driven by a special cytoplasmic dynein that is proposed to recycle the anterograde motors, IFT particles, and breakdown products of ciliary turnover back to the cell body (Pazour et al., 1998, 1999;

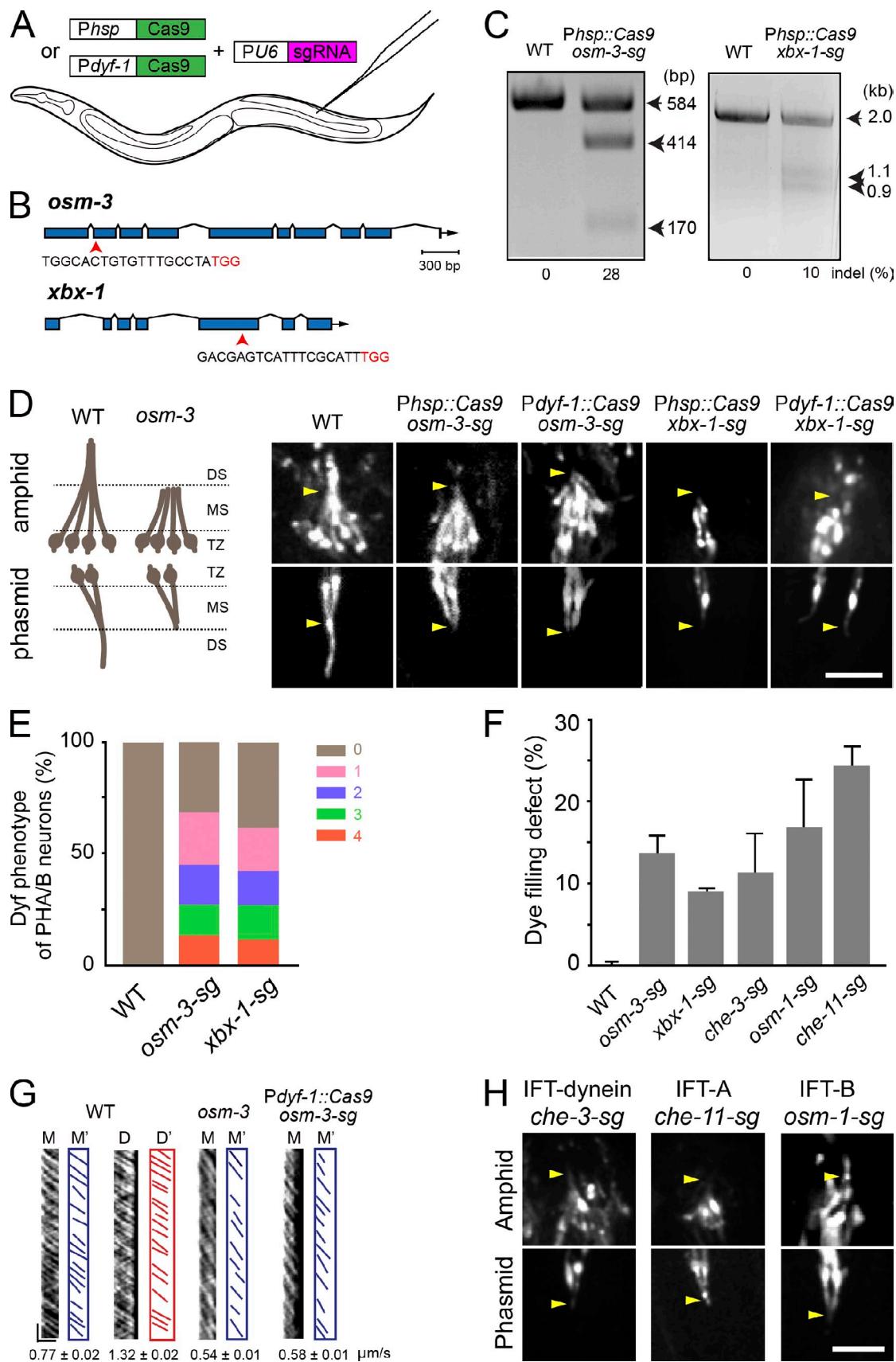
Porter et al., 1999; Signor et al., 1999; Rosenbaum and Witman, 2002; Scholey, 2003).

Cytoplasmic dyneins include cytoplasmic dynein 1 (DYNC1) and the IFT-specific cytoplasmic dynein 2 (DYNC2; Pfister et al., 2005). DYNC1 consists of a heavy chain homodimer, intermediate chains, light intermediate chains, and light chain dimers of the LC8, Tctex1, and Roadblock protein families (Kardon and Vale, 2009). These accessory subunits contribute to the structural integrity, motor regulation, and cargo specificity of the dynein complex. Moreover, cytoplasmic dynein motor activity appears to be governed by the Lis1–NudEL complex and the dynactin complex (Kardon and Vale, 2009; Roberts et al., 2013). In contrast, the composition and regulation of IFT-dynein remain unclear. Biochemical studies in *C. reinhardtii* revealed that IFT-dynein contains the heavy chain DHC1b, intermediate chains FAP133 and FAP163, light intermediate chain D1bLIC, and light chain LC8 (Pazour et al., 1998, 1999; Porter et al., 1999; Hou et al., 2004; Rompolas

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Abbreviations used in this paper: Cas, CRISPR-associated; CRISPR, clustered regularly interspaced short palindromic repeats; Dif, abnormal dye filling; IFT, intraflagellar transport; sgRNA, synthetic guide RNA; T7E1, T7 endonuclease I; WT, wild type.



**Figure 1. The somatic CRISPR-Cas9 technique generates conditional mutations in IFT components in *C. elegans* ciliated sensory neurons.** (A) Schematic illustrating the use of somatic CRISPR-Cas9 for creating conditional mutants. (B) *osm-3* and *xbx-1* gene models. Exons are in blue, and arrows indicate sgRNA sequences corresponding to exons [Table S4]. The NGG (PAM) sequences are denoted in red. (C) Representative gels showing the T7EI assay results for *osm-3* and *xbx-1*. Indels are indicated at the bottom. (D, left) Schematic of cilia in WT and *osm-3* animals. DS, distal segments; MS, middle segments;

et al., 2007; Patel-King et al., 2013). A recent study using a temperature-sensitive allele of DHC1b demonstrated that retrograde IFT regulates flagellar assembly and function but not maintenance (Engel et al., 2012). Genetic research in *Caenorhabditis elegans* revealed that mutations of *che-3* (DHC1b homologue) or *xbx-1* (D1bLIC) caused defects in retrograde IFT (Signor et al., 1999; Schafer et al., 2003).

Identification of IFT–dynein components largely relied on biochemical fractionation and proteomic analysis of flagellar proteins in *C. reinhardtii*, and this strategy does not differentiate between the integral components and the cargo molecules of IFT–dynein, as axonemal dynein complexes share subunits with IFT and cytoplasmic dyneins (e.g., LC8; Pazour et al., 1998). Moreover, mass spectrometry may not be sensitive enough to detect IFT regulators present at substoichiometric concentrations. The *C. elegans* neuronal cilia provide a complementary system to study IFT–dynein using genetic and imaging approaches. However, the efficiency of RNAi in knocking down gene activity in *C. elegans* ciliated neurons has been very limited (Pastrana, 2010). Moreover, the use of RNAi may lead to variable penetrance of the phenotype due to the incompleteness of protein depletion, and may also be accompanied by off-target effects (Pastrana, 2010). Thus, new techniques that are efficient and robust are needed to access the role of the target gene in cilia.

The clustered regularly interspaced short palindromic repeats (CRISPR)/CRISPR-associated (Cas) system has been recently demonstrated as a powerful tool for genome editing (for review see Hsu et al., 2014). The specificity of Cas9 nuclease is determined by a synthetic guide RNA (sgRNA) that base-pairs with a G(N)19NGG “protospacer” DNA sequence in the genome. Cas9 generates double-strand breaks, which are often inaccurately repaired by the nonhomologous end joining pathway, causing frameshift insertions or deletions (indels) in the target locus. We have developed the somatic CRISPR–Cas9 technique in which CRISPR–Cas9 is expressed under the control of an inducible or tissue-specific promoter to generate conditional mutants in *C. elegans* (Shen et al., 2014). Here, we first demonstrate that somatic CRISPR–Cas9 efficiently creates conditional mutations in ciliated neurons. Using this platform, we bypass the embryonic requirement of dynein-associated chains and investigate their roles in the construction of *C. elegans* neuronal cilia.

## Results and discussion

### Generation of conditional mutants in ciliated neurons of *C. elegans*

We generated transgenic animals that specifically express Cas9 under the control of either a heat-shock-inducible promoter (*Phsp-16.2* or *Phsp*) or a ciliated neuron-specific promoter

(*Pdyf-1*) and that ubiquitously express the sgRNA under the control of the *U6* gene promoter (*PU6*; Fig. 1 A; Ou et al., 2005; Shen et al., 2014). We first designed sgRNA that was complementary to the coding sequence of the *osm-3* gene or the *xbx-1* gene (Fig. 1 B). *osm-3* encodes a homodimeric kinesin-2 that is essential for building the ciliary distal segment (Snow et al., 2004), and *xbx-1* encodes an IFT–dynein light intermediate chain (Schafer et al., 2003). We chose to target *osm-3* and *xbx-1* on the basis of their easily identifiable defects in the cilium structure, IFT, and uptake of a fluorescent dye (abnormal dye filling [Dyf] phenotype). To achieve temporally controlled mutation, the expression of Cas9 was controlled by *Phsp* (Fig. 1, B and C). We transformed a plasmid containing Cas9 and sgRNA into *C. elegans*, and in the resultant transgenic animals, we refer to the sgRNA targeting *osm-3* or *xbx-1* as *osm-3-sg* or *xbx-1-sg*, respectively. Because T7 endonuclease I (T7EI) recognizes and cleaves the imperfectly matched DNA (Hsu et al., 2014), we applied the T7EI assay to detect the indels in *osm-3-sg* and *xbx-1-sg* animals whose embryos were treated with heat shock. We first PCR-amplified the genomic DNA fragments containing the target sites from these transgenic animals. After T7EI digestion, these PCR fragments were cut into two small fragments of the expected sizes; however, the fragments were intact in wild-type (WT) animals (Fig. 1 C), which indicates that somatic CRISPR–Cas9 induced molecular lesions in the *osm-3* and *xbx-1* loci.

We next used a compound microscope with a 100× objective lens to examine the Dyf phenotype (Fig. S1). By examining four phasmid ciliated neurons (PHA/B/L/R) of *osm-3-sg* and *xbx-1-sg* conditional mutants, we found that 69% and 62% of the animals showed defects in Dil uptake, respectively ( $n = 51$  and 26; Figs. 1 E and S1 A). All neurons showed complete uptake of Dil in 100% of WT animals ( $n > 100$ ; Fig. 1, E and F). We noticed that the mosaic Dyf phenotype ranged from 14% to 24% for *osm-3-sg* and from 12% to 19% for *xbx-1-sg* in one to four phasmid neurons (Fig. 1 E), which is a common feature in conditional mutants. We also assessed the Dyf phenotype using a fluorescence stereoscope and detected the lower Dyf penetrance (Fig. 1 F). This result may have occurred because the stereoscope is sensitive enough to detect fluorescence from a single neuron, and we only scored the Dyf phenotype in animals with phasmid neurons that completely failed to take up Dil (Figs. 1 E and S1, A and B).

To induce spatially controlled mutations, we used *Pdyf-1* to express Cas9. Under a fluorescence stereoscope, we found that 92% ( $n = 50$ ) and 56% ( $n = 27$ ) of transgenic animals expressing *Pdyf-1::Cas9*; *osm-3-sg* and *Pdyf-1::Cas9*; *xbx-1-sg* showed the Dyf phenotype, respectively. Thus, the expression of Cas9 under the control of a heat-shock promoter or a ciliary-specific promoter allows efficient creation of conditional mutants

TZ, transition zone. (D, right) The morphology of amphid (top) and phasmid (bottom) cilia visualized using OSM-6::GFP. Arrowheads indicate junctions between the middle and distal segments. The length of the cilia are  $8.2 \pm 0.1$  μm (amphids [A]) or  $7.9 \pm 0.3$  μm (phasmids [P]) in WT,  $4.8 \pm 0.1$  μm (A) or  $3.9 \pm 0.2$  (P) μm in *Phsp-16.2::Cas9*; *osm-3-sg*, and  $4.5 \pm 0.1$  μm (A) or  $3.5 \pm 0.1$  μm (P) in *Pdyf-1::Cas9*; *osm-3-sg*.  $n = 23$ –81. Mean  $\pm$  SE. Bar, 5 μm. (E) Dye-filling defects under a 100× objective lens. 0–4 and their respective colors represent the number of PHA/B neurons defective in dye-filling assays.  $n = 26$ –51. (F) Dye-filling defects under a fluorescence stereoscope.  $n = 150$ –315. Mean  $\pm$  SE (error bars). (G) Kymographs (M or D) and corresponding lines (M' or D') of OSM-6::GFP motility. Error bars indicate mean  $\pm$  SE. Horizontal bar, 2 μm; vertical bar, 5 s.  $n = 120$ –323. (H) Cilium morphology in conditional mutants visualized by OSM-6::GFP. Arrowheads indicate the positions where the junctions between the middle and distal segments should be in WT animals. Bar, 5 μm.

of *osm-3* and *xbx-1*. By comparing mutation efficacies and the penetrance of ciliary phenotypes between *osm-3* and *xbx-1*, we noticed that the higher indel frequency (28% for *osm-3*, 10% for *xbx-1*) may correlate with the higher dye filling defect (15% for *osm-3*, 9% *xbx-1*; Fig. 1, C and F).

Next, we examined the defects in cilium structure and IFT in *osm-3*-*sg* and *xbx-1*-*sg* animals using the OSM-6/IFT52::GFP reporter, which undergoes biphasic IFT along the entire cilia (Fig. 1 D). The *osm-3*-*sg* animals lost their ciliary distal segments, and the cilium length was reduced from 8 μm to 4–5 μm (Fig. 1 D), phenocopying *osm-3(p802)* null mutants (Snow et al., 2004). We further performed live imaging analysis of IFT in *osm-3*-*sg* mutants and found that the IFT velocity was reduced from  $0.77 \pm 0.02$  μm/s (mean ± SE,  $n = 213$ ) in the WT animals to  $0.58 \pm 0.01$  μm/s ( $n = 323$ ) in the remaining middle segments, which is also comparable to the IFT speed in *osm-3(p802)* mutants (Fig. 1 G). In *xbx-1*-*sg* conditional mutants, OSM-6::GFP formed aggregates around the transition zone and no IFT was detected in the residual cilia (Fig. 1 D), which are similar to those observed in *xbx-1(ok279)* deletion mutants (Schafer et al., 2003). To extend our initial successes to other IFT components, we conditionally mutated the IFT-dynein heavy chain (*che-3*), IFT particle A (*che-11*), and IFT particle B (*osm-1*) components by using the heat-shock strategy (Signor et al., 1999; Scholey, 2003). We showed that these conditional mutants reproduced the defects in cilium structure caused by their corresponding null alleles, and that the Dyf penetrance ranged from 10% to 25% in four plasmid ciliated neurons (Fig. 1, F and H). Collectively, somatic CRISPR–Cas9 can be used to generate conditional mutations of IFT motors (*osm-3*, *xbx-1*, and *che-3*) and IFT particles (*che-11* and *osm-1*).

#### The functions of cytoplasmic dynein components in ciliogenesis

The *C. elegans* genome encodes 14 dynein-associated chains (Hao et al., 2011), five of which are essential for embryonic development (WormBase), including the intermediate chain (DYCI-1), light intermediate chain (DLI-1), light chain LC8 (DLC-1), Tctex-type light chain (DYL-T-3), and roadblock-type light chain (DYRB-1; Fig. 2 A). We first generated transgenic animals that expressed Cas9 under the control of the *Phsp* promoter and ubiquitously expressed sgRNAs that target one site for *dylt-3* and *dyrb-1* or two sites for *dyci-1*, *dlc-1*, and *dli-1* (Fig. 2 A). Our T7EI assays detected the expected molecular lesions (Fig. 2 B). Although the heat-shock treatment did not alter the embryonic viability of WT embryos, the conditional mutant embryos exhibited embryonic lethality with penetrances ranging from 40% to 55% after heat shock (Fig. 2 C). These data indicated that we generated conditional mutants of embryonically essential dynein subunits.

Next, we used these conditional mutants to study the contribution of the dynein components to cilium formation. We first showed that 29% (*dyci-1*), 19% (*dlc-1*), and 18% (*dli-1*) of the conditional mutants failed to take up DiI, whereas <4% of *dylt-3* or none of *dyrb-1* conditional mutants developed the Dyf phenotype (Fig. 3 A). Consistently, we found that the cilium length was significantly reduced from 8.3 μm in WT animals to 4.4 μm (*dyci-1*), 4.9 μm (*dlc-1*), and 4.8 μm (*dli-1*) in the conditional

mutants but was not altered in *dylt-3*-*sg* or *dyrb-1*-*sg* animals (Fig. 3, A and B). IFT cargo includes IFT particles and ciliary receptors such as a transient receptor potential vanilloid channel OSM-9 (Qin et al., 2005). We did not find that OSM-6::GFP or OSM-9::GFP moved in the remaining cilia of the *dyci-1*-*sg* and *dlc-1*-*sg* conditional mutants (Figs. 3 B and S3 A). These ciliary defects are similar to those detected in *che-3*- and *xbx-1*-null mutants. To further test whether DYCI-1 and DLC-1 function with IFT-dynein, we determined the cellular localization of GFP-tagged DYCI-1 and DLC-1. We showed that they were distributed along the cilia and underwent the same biphasic IFT as XBX-1 (Fig. 3 C and see Fig. 5 B). Furthermore, we genetically introduced XBX-1::YFP into the *dyci-1*-*sg* conditional mutant and GFP::DYCI-1 into the *xbx-1(ok279)* mutant. We found that XBX-1::YFP and GFP::DYCI-1 formed aggregates around the transitional zone and were not motile in cilia (Fig. 3, D and E). The interdependence of XBX-1 and DYCI-1 in IFT further indicated that they may function together with IFT-dynein.

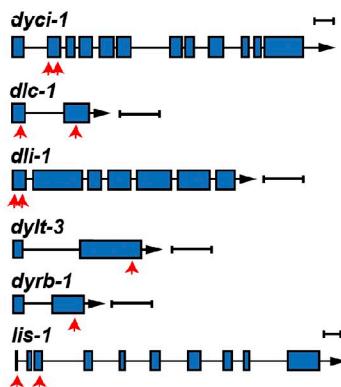
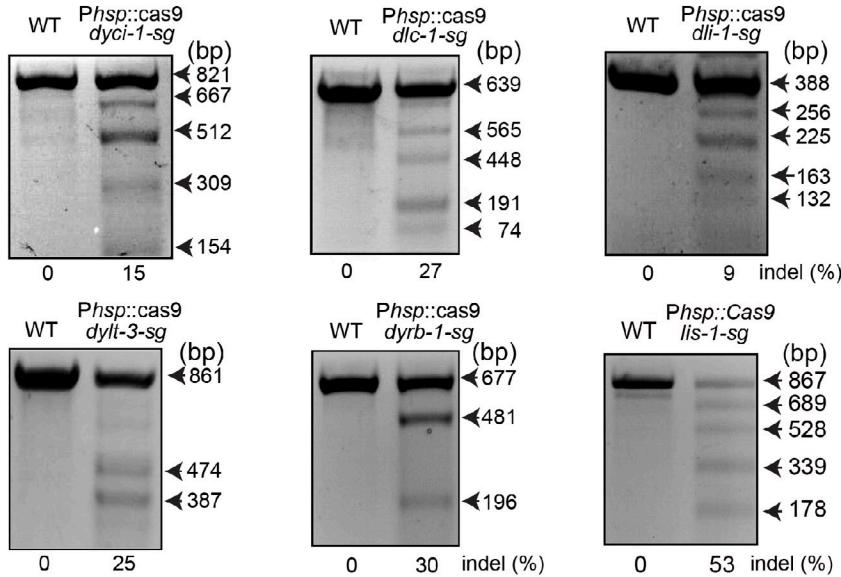
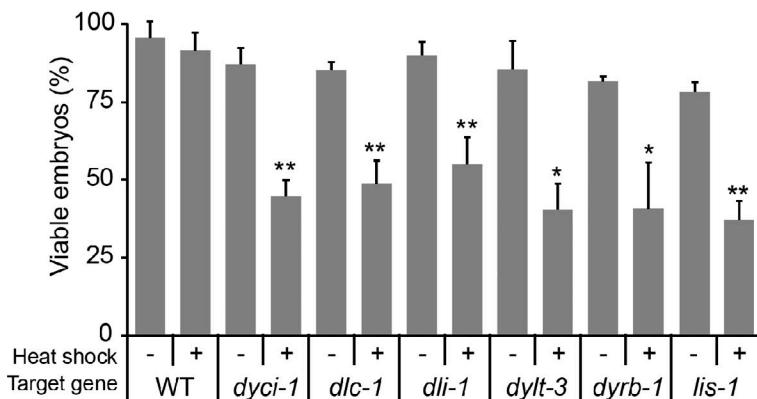
We studied how DLI-1 contributes to ciliogenesis. We did not find that OSM-6::GFP formed aggregates within the cilia of *dli-1*-*sg* conditional mutants (Fig. 3 B), nor did we detect the ciliary localization of DLI-1::GFP in transgenic animals expressing the *Pdli-1::dli-1*::GFP or *Pdyf-1::gfp::dli-1* reporter (Figs. 3 C and S2 A). Instead, we found that DLI-1::GFP was distributed along the dendrite in these neurons (Fig. S2 B) and that OSM-6::GFP formed aggregates in the dendrites of *dli-1*-*sg* conditional mutants (Fig. S2 C). To further examine the function of DLI-1 in the retrograde dendritic transport, we compared the transport velocities of OSM-6::GFP in the dendrites of WT and *dli-1* conditional mutants. Consistent with the previous measurements (Signor et al., 1999), OSM-6::GFP moved along the dendrite at the anterograde speed of  $0.75 \pm 0.05$  μm/s ( $n = 33$ ) and at  $0.90 \pm 0.04$  μm/s ( $n = 30$ ) in WT animals. However, we did not detect the retrograde movement of OSM-6::GFP in the dendrites of the *dli-1* conditional mutants, whereas we found that OSM-6::GFP moved at the anterograde speed of  $0.68 \pm 0.02$  μm/s ( $n = 33$ ) in these dendrites. These results indicate that DLI-1 indirectly contributes to cilium formation by regulating the retrograde dendritic transport.

#### Dynein subunits undergo diverse IFT

To understand the transport behavior of IFT-dynein subunits, we generated kymographs from time-lapse movies of IFT of GFP-tagged dynein subunits (Videos 1–4). We first quantified the transport frequency as the number of transport events per minute. OSM-6::GFP and XBX-1::YFP both showed a frequency of 16 events/min; however, DYCI-1, DLC-1, and DYL-T-3 displayed frequencies of ~3, 9, and 6 events/min, respectively (Fig. 4 A). We next examined their turnaround sites. The majority of OSM-6 and XBX-1 turned around at the distal tip, whereas DYCI-1, DLC-1, and DYL-T-3 returned at the middle and distal segments (Fig. 4, B and C). 76% of the GFP::DYCI-1 puncta returned along the distal segment and very few reached the distal tip, and 24% of DYCI-1 turned around at the tip of the middle segment. 60% of the DLC-1::GFP puncta turned back at the distal segment, whereas 40% of DLC-1 returned at the middle tip. Finally, 69% of the DYL-T-3::GFP puncta returned

**A**

Essential Dynein Subunits across Species			
Dynein subunits family	C. elegans	H. sapiens	C. reinhardtii
Heavy chain	che-3	DYNC2H1/DHC2	DHC1b
Intermediate chain	dyci-1	DYNC1I1/2	D1bLIC/FAP133
Light intermediate chain	dli-1 xbx-1	DYNC1I1/2 DYNCL2L1/D2LIC	D1bLIC
Light chain	dlc-1	DYNLL1/2	LC8/FLA14
Light chain Tctex	dylt-3	DYNLT1/3 DYNLL1/2	Tctex1
Light chain roadblock	dyrb-1	DYNLRB2	DLC7a/b

**B****C**

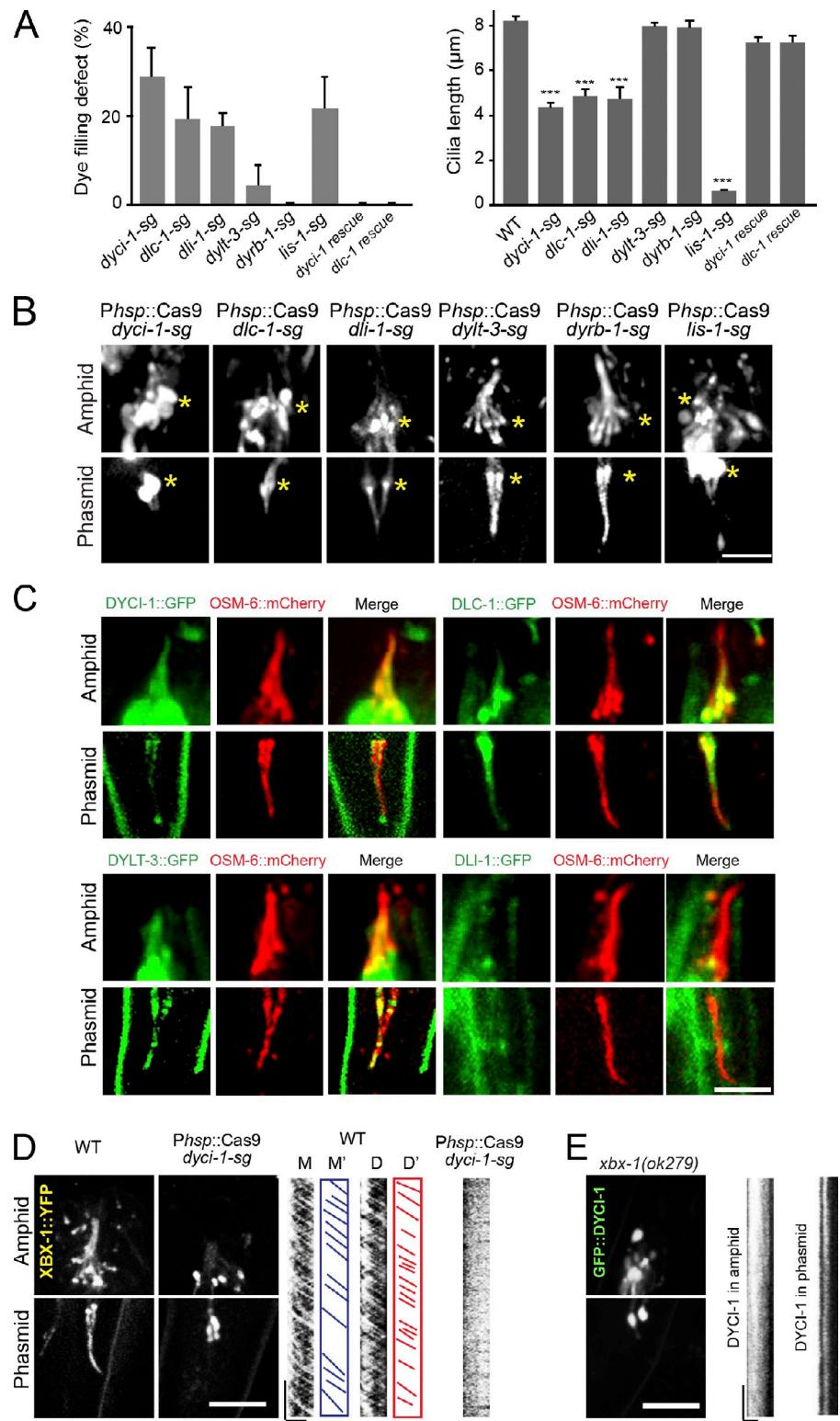
in the middle segment, not even reaching the middle tip (Fig. 4, B and C). These data suggest that multiple retrograde IFT pathways may operate within a single cilium.

#### Lissencephaly-1 regulates retrograde IFT

We studied the function of lissencephaly-1 in cilium formation. Lis1 is thought to function as a “clutch” to suppress cytoplasmic dynein motility and to cause the dynein to tightly bind microtubules (Huang et al., 2012). We generated *lis-1-sg* conditional mutants that exhibited the expected molecular lesions and embryonic

**Figure 2. Conditional mutations in six embryonically essential dynein components.** (A, left) The essential dynein subunits and known IFT-dynein components across species. (A, right) Gene models of *dyci-1*, *dlc-1*, *dli-1*, *dylt-3*, *dyrb-1*, and *lis-1*. Exons are in blue and red arrows indicate one or two sgRNA sequences targeting dynein components (Table S4). Bars, 300 bp. (B) Representative gels showing the results of T7EI assays. One target site generated two small fragments. (C) Embryonic lethality was determined by quantifying viable embryos after heat-shock treatment.  $n = 97\text{--}242$ ; mean  $\pm$  SE (error bars); \*,  $P < 0.05$ ; \*\*,  $P < 0.01$ .

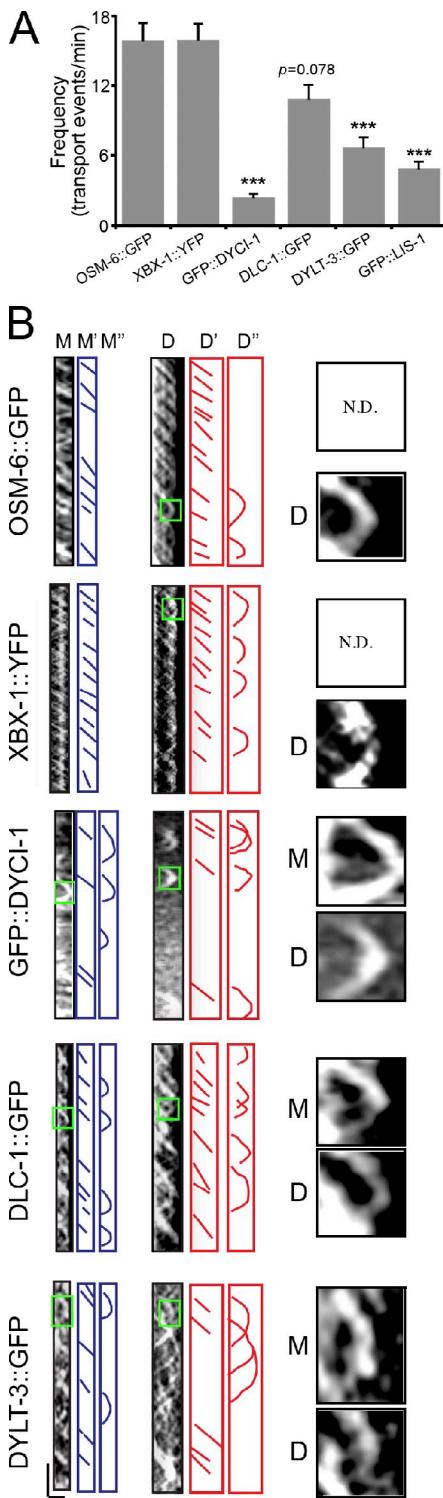
**Figure 3. Ciliary defects in conditional mutants of embryonically essential dynein components.** (A) Defects of dye-filling assays (left) and the cilium length (right). Mean  $\pm$  SE (error bars);  $n = 34\text{--}381$  from three generations. \*\*\*,  $P < 0.001$ . (B) Amphid and phasmid cilia visualized by OSM-6::GFP. Asterisks indicate transition zones. Bar, 5  $\mu\text{m}$ . (C) Ciliary localization of DYCI-1, DLC-1, DYLT-3, and DLI-1 (GFP) and OSM-6::mCherry (red). (D and E) Localization (left) and motility (kymographs, right) of XBX-1::YFP in WT and *dyci-1* conditional mutants (D) or GFP::DYCI-1 in *xbx-1* mutants (E). Micrograph bar, 5  $\mu\text{m}$ ; kymograph horizontal bar, 2  $\mu\text{m}$ ; vertical bar, 5 s.



In conclusion, this study demonstrated that embryonically essential dynein-associated chains behave distinctly in *C. elegans* neuronal cilia: the intermediate chain DYCI-1, the light chain LC8/DLC-1, and the dynein regulator LIS-1 undergo biphasic IFT and are essential for retrograde IFT; the light intermediate chain DLI-1 regulates the dendritic transport of IFT particles; the light chain Tctex-type DYLT-3 and roadblock-type

DYRB-1 are dispensable for ciliogenesis (Fig. 5, A and B). These findings suggest that cytoplasmic dynein and IFT–dynein have unique compositions but share essential components and regulatory mechanisms.

To visualize IFT of dynein subunits, this work relied on the GFP fusion technique, a standard protocol that has generated reliable results including ones related to IFT–dynein



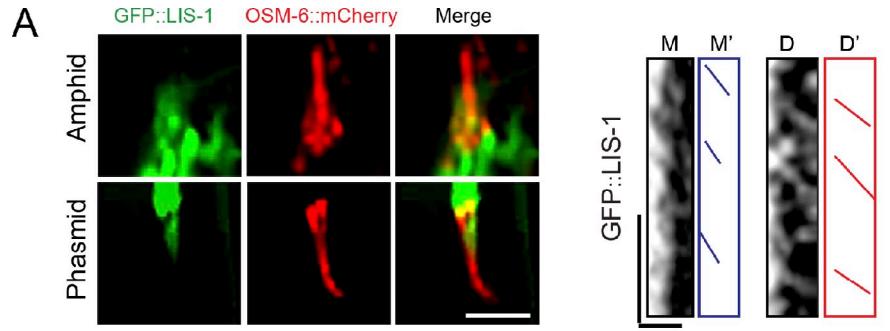
**Figure 4. Transport behaviors of dynein subunits in cilia.** (A) IFT frequencies of IFT particle and dynein subunits.  $n = 64\text{--}212$ ; mean  $\pm$  SE (error bars). \*\*\*,  $P \leq 0.001$ . (B) Kymographs (left) of DYCI-1, DLC-1, DYLT-3, and XBX-1 in the middle (M) or distal segments (D). M'' and D'' indicate turnaround events. M and D (right) are 4x enlarged images of turnarounds in the green boxes on the left. Horizontal bar, 2  $\mu\text{m}$ ; vertical bar, 5 s. (C) Distribution of turnaround events along the cilia. The y axis indicates the percentage of turnaround events at the specific region of cilia among the total turnaround events.  $n = 58\text{--}140$ . The broken line shows the junction between the middle and distal segments.

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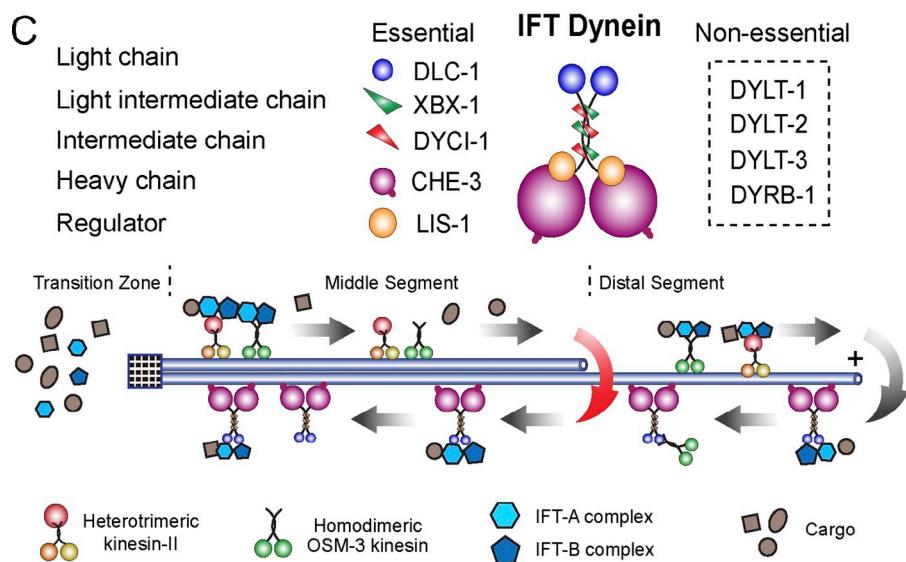
(Mello et al., 1991; Signor et al., 1999; Schafer et al., 2003). Despite the fact that we injected DNA with a low concentration ( $\sim 10 \text{ ng}/\mu\text{l}$ ), below which the GFP fluorescence is invisible, and that we did not notice any ciliary phenotype in these animals, this method can cause the overexpression of the transgene. Dynein is a holoenzyme complex with a strict stoichiometry, and the overexpression of subunits or GFP tagging could result in artifacts because some complexes would be disrupted or not fully assembled. We thus performed the

rescue experiments to examine whether the overexpression of GFP-tagged dynein subunits could rescue ciliary phenotypes in the corresponding conditional mutants. We studied *dyci-1* and *dlc-1* as they regulate ciliogenesis. In these experiments, we first made all the possible synonymous mutations in the target sites of transgenes such that they would not be cleaved by CRISPR–Cas9 (Fig. S3 B). We transformed GFP-tagged *mdyci-1* (m for synonymous mutation) and *mdlc-1* into conditional mutants, and we did not detect any Dyf phenotype after

**Figure 5. IFT of LIS-1, IFT velocities, and the proposed model.** (A) Colocalization of GFP::LIS-1 with OSM-6::mCherry (red) in cilia (left), and a kymograph of GFP::LIS-1 motility (right). Micrograph bar, 5  $\mu$ m; kymograph horizontal bar, 2  $\mu$ m; vertical bar, 5 s. (B) Velocities of IFT particle and dynein subunits (mean  $\pm$  SE). (C) A proposed model of subunits in the IFT-dynein complex and multiple turnaround sites on the cilium.



Proteins	Anterograde ( $\mu$ m/s)				Retrograde ( $\mu$ m/s)				n
	Middle segment	n	Distal segment	n	-	+	n	n	
OSM-6::GFP	0.77 $\pm$ 0.02	213	1.32 $\pm$ 0.02	231	-1.19 $\pm$ 0.03	0.03	118		
DLC-1::GFP	0.81 $\pm$ 0.02	110	1.27 $\pm$ 0.04	152	-1.13 $\pm$ 0.05	0.05	48		
DYCI-1::GFP	0.77 $\pm$ 0.06	59	1.08 $\pm$ 0.05	70	-1.10 $\pm$ 0.08	0.08	29		
DYLT-3::GFP	0.76 $\pm$ 0.05	68	1.18 $\pm$ 0.07	45	-1.07 $\pm$ 0.05	0.05	78		
XBX-1::YFP	0.87 $\pm$ 0.02	117	1.31 $\pm$ 0.03	127	-1.22 $\pm$ 0.05	0.05	38		
GFP::LIS-1	0.86 $\pm$ 0.02	109	1.13 $\pm$ 0.05	85	-1.14 $\pm$ 0.04	0.04	57		



inducing Cas9 expression by heat-shock treatment (*GFP::mdyci-1*,  $n = 59$ ; *mdlc-1::GFP*,  $n = 87$ ; Fig. 3 A). We further examined the ciliary structure and measured the cilium length in these animals, and we did not find any obvious abnormality compared with those in WT animals (Figs. 3 A and S3 C). These rescue results indicated that GFP fusion or overexpression of DYCI-1 and DLC-1 does not affect IFT-dynein's function. Because the potential off-target effect is a major concern for Cas9, our results not only demonstrated the functionality of GFP fusion, but also alleviated the concern of off-target cleavages.

The GFP-tagged dynein subunits undergo IFT at the same speeds as other IFT motors (Fig. 5 B), which suggests that they are probably associating with the dynein complex. We did not detect DLI-1::GFP in the cilia using the same expression strategy, suggesting that the ciliary location of GFP-tagged dynein

subunits can be specific (Fig. 3 C). It remains unknown why DLI-1 only functions within the dendrites but does not enter cilia. The emerging evidence suggests that a selective ciliary gate may regulate protein trafficking between cilia and the general cytoplasm (Ishikawa and Marshall, 2011), and DLI-1 may not be able to pass such a gate at the base of cilia.

Our results with IFT-dynein are consistent with biochemical data from other organisms, and also provide new insights. For example, LC8/DLC-1 was proposed as an essential component for retrograde IFT from *C. reinhardtii* (Pazour et al., 1998). However, LC8 can be the subunit or cargo—or both—of IFT-dynein, as it is a known component in axonemal dyneins. Our study from nonmotile cilia enables us to conclude that LC8 can be an IFT-dynein subunit. Similarly, a recent study used the biochemical approach to study IFT-dynein composition in vertebrates, and their data are also largely complementary to ours

(Asante et al., 2014). However, Lis1 was not identified in IFT-dynein fraction but was localized in cilia by immunofluorescence (Asante et al., 2014). We found that GFP::LIS-1 was not associated with IFT-dynein all the time (Fig. 5 A), and the biochemical fractionation must be difficult to detect the transient interaction. Our data also indicate that DYCI-1 is an integral IFT-dynein component. The previous work using a transcriptional GFP reporter did not detect the expression of *dyci-1* in *C. elegans* ciliated neurons (Hao et al., 2011). By increasing the length of the 5' UTR regulatory sequence, we detected the expression of *Pdyci-1::gfp* in ciliated neurons (Fig. 3 C). Consistently, biochemical studies identified FAP133/DYCI-1 as an IFT-dynein component (Rompolas et al., 2007), and RNAi experiments in *Trypanosome brucei* showed that DIC5/DYCI-1 regulates retrograde IFT (Blisnick et al., 2014). Moreover, we showed that DYLT-3 and DYRB-1 do not regulate ciliogenesis (Fig. 3, A and B). Dynein-associated chains may complement each other in cilium formation, and the generation of double or multiple conditional mutations by somatic CRISPR–Cas9 may dissect their redundant roles in ciliogenesis. Interestingly, the single or double genetic deletions of *C. elegans* Tctex-type light chain DYLT-1 or DYLT-2 do not cause defects in cilium assembly, even though they undergo biphasic IFT (Hao et al., 2011). We cannot exclude the possibility that these dynein light chains regulate ciliogenesis, but they may also likely modulate currently unidentified physiological events in cilia.

Lis1 has been found to induce slow or nonmoving dynein (Huang et al., 2012). We showed that GFP::LIS-1 is distributed in the middle segment (Fig. 5 A), where it may inactivate IFT-dynein during anterograde IFT. We observed biphasic but infrequent IFT of LIS-1 in *C. elegans* (Fig. 5, A and B), which is consistent with the brief motility of Lis1 in *Aspergillus nidulans* (Egan et al., 2012). Because limited GFP::LIS-1 enters the ciliary distal segment where retrograde IFT commences, IFT-dynein motility may also require the detachment of LIS-1, and a substoichiometric amount of LIS-1 at the distal tip may initiate IFT. In Lis1 mutants, cytoplasmic dynein and its cargo move at normal speeds with reduced frequencies in *A. nidulans* (Egan et al., 2012; Huang et al., 2012). However, we did not observe retrograde IFT in the *lis-1::sg* conditional mutants, which suggests that LIS-1 may be more critical for the motility of IFT-dynein. Studies from mammalian motile cilia and *C. reinhardtii* flagella reported that Lis1 regulates the activity of the outer dynein arm (Pedersen et al., 2007), indicating that Lis1 plays manifold roles in the regulation of cytoplasmic dynein, IFT-dynein, and axonemal dynein.

In summary, the somatic CRISPR–Cas9 technique enables the functional analysis of embryonically essential dynein subunits in *C. elegans* ciliogenesis. These dynein-associated molecules may regulate the motor processivity or function as a switch to change the direction of motion, and single molecule biophysics studies should be pursued in the future to better understand the function of these dynein-associated chains. The present study focuses on the ciliary roles of embryonically essential dynein components, and these conditional mutants are also valuable resources for the study of other dynein-based processes. This

method is also amenable to investigations of the ciliary function of other essential embryonic genes as well as the formation and function of other organelles.

## Materials and methods

### *C. elegans* strains, genetics, and DNA manipulations

*C. elegans* strains were raised on nematode growth media (NGM) plates seeded with the *Escherichia coli* strain OP50 at 20°C using standard methods. CRISPR–Cas9 constructs were produced according to a previously described method (Shen et al., 2014). In brief, to generate the conditional knockout vectors, CRISPR–Cas9 target sequences were added to CRISPR–Cas9 vectors by PCR using Phusion high-fidelity DNA polymerase (New England Biolabs, Inc.) with the primers listed in Table S3 followed by self-ligation and transformation into *E. coli*. GFP-tagged dyneins were constructed by insertion of genomic sequences (3 kb promoter plus coding region) into pPD95.77 (containing GFP and unc-54 3' UTR) for endogenous expression or into POG1264 (containing *dyf-1* promoter, GFP, and unc-54 3' UTR) for ciliated neuron specific expression, except that DYLT-3::GFP fusion was generated by PCR gene splicing by overlap extension (SOEing) of a *dylt-3* genomic sequence (3 kb promoter plus coding region) with GFP::unc-54 3' UTR DNA fragments. Transgenic *C. elegans* were generated by the germ line microinjection of DNA plasmids at 10–50 ng/μl and a selection marker pRF4[rol-6(su1006)], *Podr-1::dsRed*, WT *unc-76* gene or *Pegl-17::Myri-mCherry + Pegl-17::mCherry::his-24* into *N<sub>2</sub>* or *unc-76(e911)* hermaphrodites. The strains, primers, and plasmids are listed in Tables S1–S3.

### Molecular assays

Molecular lesions induced by CRISPR–Cas9 were detected according to previously described methods (Shen et al., 2014). In brief, worms were lysed with 10x PCR buffer ( $Mg^{2+}$ ) and were used as templates for amplifying DNA fragments containing CRISPR–Cas9 targets. PCR products were purified by QIAquick PCR Purification kit (QIAGEN) and digested by T7EI at 37°C for 30 min. The indel percentage was determined using the formula  $100 \times \{1 - [1 - (b + c)/(a + b + c)]^{1/2}\}$ , where  $a$  is the integrated intensity of the intact band and  $b$  and  $c$  are the integrated intensities of the digested small bands.

### Heat-shock treatment, dye-filling assay, and viability assay

Heat-shock treatment was performed at 33°C for 1 h at 8 h after egg laying. The dye-filling assay was performed according to a previously described protocol (Snow et al., 2004). In brief, young adult worms were collected into ~200 μl M9 solution, and mixed with ~200 μl dyes (DiI, 1,1'-dioctadecyl-3,3',3',3'-tetramethylindo-carbocyanine perchlorate, or DiO, 3,3'-dioctadecyloxacarbocyanine perchlorate) at a working concentration (20 μg/ml). After incubation at room temperature in a dark place for ~30–60 min, animals were transferred to a seeded NGM plate, and examined for dye uptake 1 h later. For the viability assay, the number of viable adult worms was divided by the number of originally laid eggs.

### Live cell imaging

*C. elegans* were anesthetized with 0.1 mmol/liter levamisole in M9 buffer, mounted on 2% agar pads, and maintained at 22°C. Our imaging system includes an Axio Observer Z1 microscope (Carl Zeiss) equipped with a 100x, 1.45 NA objective lens, an EM CCD camera (iXon+ DU-897D-C00-#BV-500; Andor Technology), and the 488 nm and 568 nm lines of a Sapphire CW CDRH USB Laser System (Coherent) with a spinning disk confocal scan head (CSU-X1 Spinning Disk Unit; Yokogawa Electric Corporation). Time-lapse images were acquired with an exposure time of 200 ms by μManager ([www.micro-manager.org](http://www.micro-manager.org)). We used ImageJ software (<http://rsbweb.nih.gov/ij/>) to process the images.

### Statistical analysis

Student's *t* test and  $\chi^2$  analysis were used to examine significant differences in ciliary phenotypes between WT and mutant animals, as indicated in the figure legends.

### Online supplemental material

Fig. S1 (related to Fig. 1), shows the Dyf phenotype in *osm-3::sg* or *xbx-1::sg* animals under a 100x objective lens and a fluorescence stereoscope, respectively. Fig. S2 (related to Fig. 3) shows DLT-1 localization and function in ciliated neurons. Fig. S3 (related to Fig. 3 and Fig. 4) indicates that GFP fusion or overexpressed DYCI-1 and DLC-1 do not affect IFT-dynein's function. Table S1 lists strains used in this study. Table S2 and S3 list all PCR products and plasmids for construction of transgenic *C. elegans*. Table S4

lists the target sequence of each gene. Table S5 gives the primers for molecular analysis. Videos 1–4 (related to Fig. 4) show the IFT movements of embryonically essential dynein subunits in cilia (GFP::DYCI-1, DLC-1::GFP, DYLT-3::GFP, and GFP::LIS-1). Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.201411041/DC1>.

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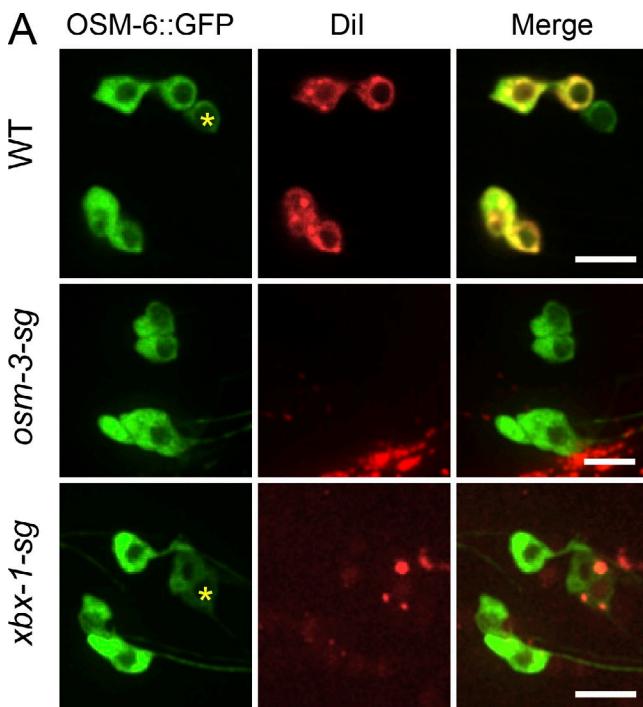
The authors declare no competing financial interests.

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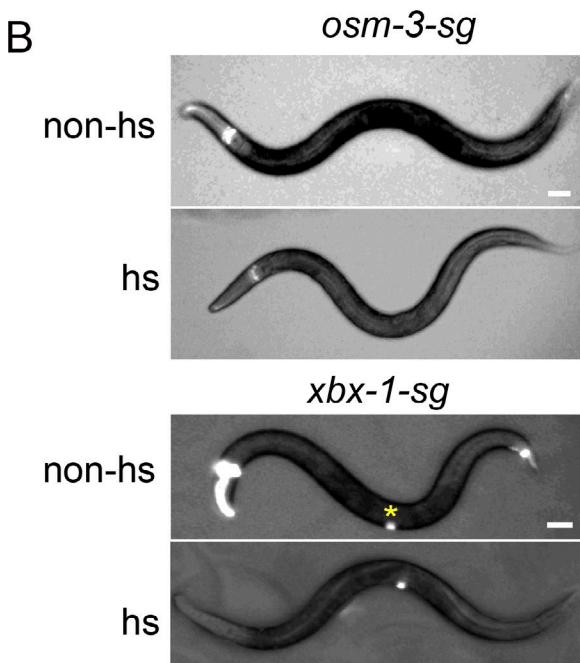
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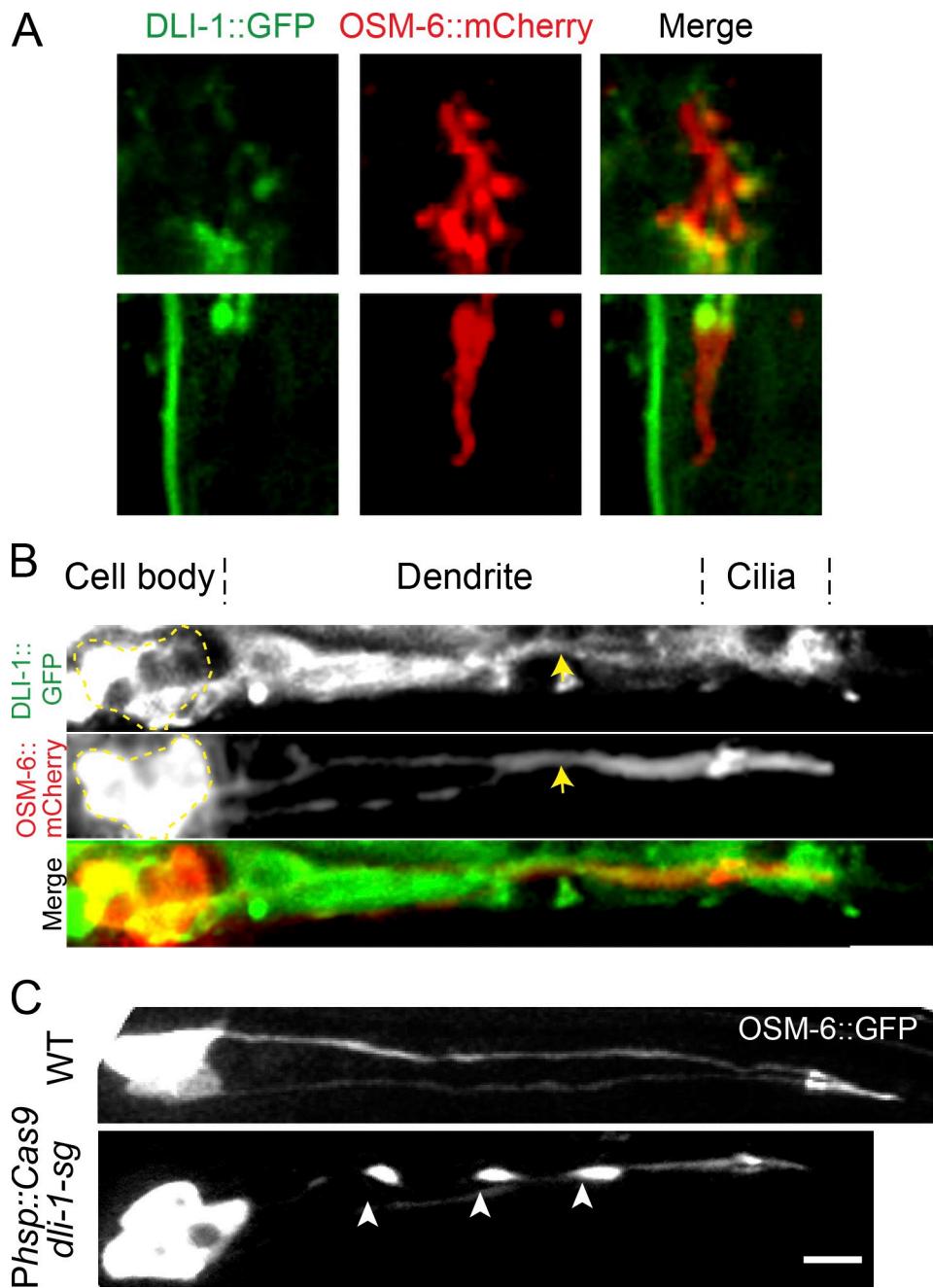
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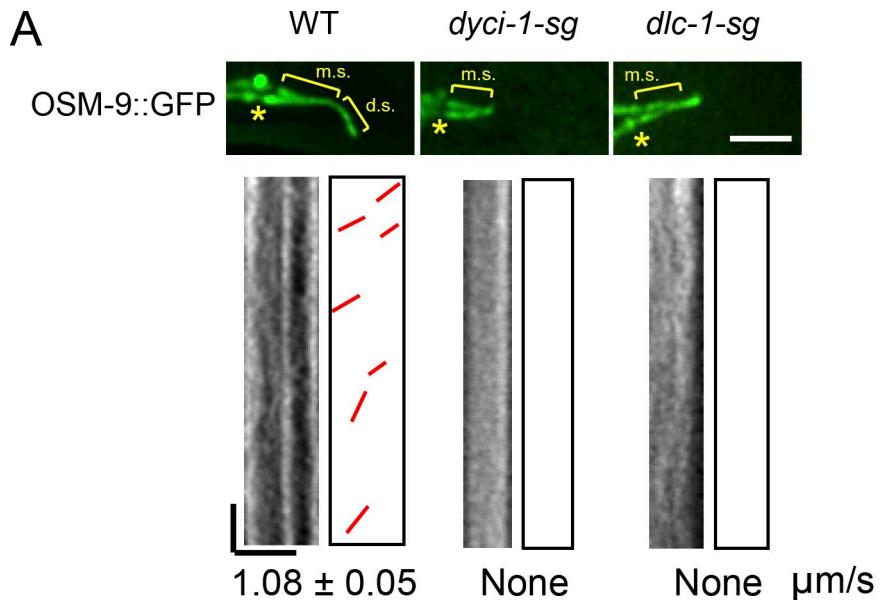
Li et al., <http://www.jcb.org/cgi/content/full/jcb.201411041/DC1>

**Figure S1. The Dyf phenotype in *osm-3-sg* or *xbx-1-sg* animals.** (A) The dye-filling phenotype of four phasmid neurons (PHA/B/L/R) of WT, *osm-3-sg*, and *xbx-1-sg* conditional mutants under a 100x objective lens. The asterisk indicates a PQR neuron, which does not take up Dil in WT animals. Bar, 5  $\mu$ m. (B) The Dyf defects in *osm-3-sg* and *xbx-1-sg* conditional mutants as observed under a fluorescence stereoscope. The head and the tail of the ciliated sensory neurons are stained by the red fluorescent Dil dye in WT animals, whereas a dye-filling defect is observed in conditional mutant animals. The asterisk indicates the transgenic marker. Non-hs, nonheat shock; hs, heat shock. Bar, 50  $\mu$ m.



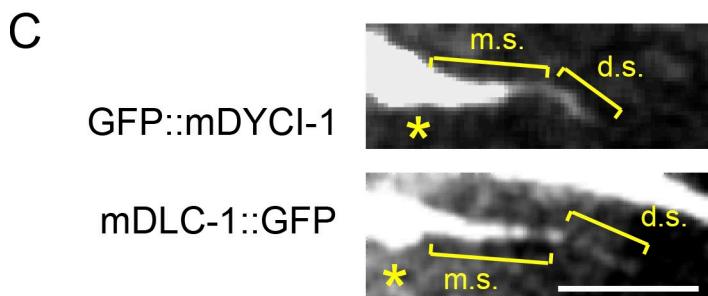


**Figure S2. DLI-1 functions in *C. elegans* ciliated neurons.** (A and B) Animals expressing *Pdli-1::dli-1::GFP* illustrate that DLI-1 does not enter the cilia (A) but localizes in the cell body and dendrites of ciliated neurons (B, arrows). The broken outlines indicate the junctions between segments within ciliated neurons. (C) Aggregates of OSM-6::GFP (arrows) along the dendrite are observed in *dli-1* conditional mutant animals. Bar, 5  $\mu$ m.



**B**

		sgRNA target	PAM
<i>dyci-1-sg</i> T1	WT	<b>aggatattctccgtcaagt</b> AGG	
	mut.	<b>aagacatatgagacagg</b> t <b>CGG</b>	
<i>dyci-1-sg</i> T2	WT	<b>agtgtacagacagcgtgca</b> CGG	
	mut.	<b>agcgtccaaaccgc</b> t <b>tgta</b> CTG	
<i>dlc-1-sg</i> T1	WT	<b>caaccctggcattgcattcg</b> t <b>CGG</b>	
	mut.	<b>ccaca</b> t <b>ggca</b> c <b>tgtat</b> AGG	
<i>dlc-1-sg</i> T2	WT	<b>gccatcgactgcgcccaccc</b> AGG	
	mut.	<b>gccatcgactgcgcccac</b> a <b>cAAG</b>	



**Figure S3. GFP-tagged DYCI-1 and DLC-1 with synonymous mutations rescued ciliary phenotypes in the corresponding conditional mutants.** (A) The retrograde movement of OSM-9::GFP in cilia of WT, *dyci-1-sg*, and *dlc-1-sg* conditional mutant animals. (top) OSM-9::GFP localization. (bottom) Kymographs and corresponding lines of the movement. Bars: (micrograph bar on top) 5  $\mu\text{m}$ ; (kymograph horizontal bar) 2  $\mu\text{m}$ ; (vertical bar) 5 s. (B) Synonymous mutations (red) of the target sites in *dyci-1* and *dlc-1*. (C) The localization of GFP-tagged DYCI-1 and DLC-1 with synonymous mutations in cilia of *dyci-1-sg* and *dlc-1-sg* conditional mutant animals. Bar, 5  $\mu\text{m}$ . m.s., middle segment; d.s., distal segment. The asterisks indicate the transition zone.

Table S1. *C. elegans* strains used in this study

Strain name	Genotype	Method
SP2101	ncl-1(e1865) unc-36(e251); osm-6(p811); mnls17[Posm-6::osm-6::GFP; unc-36(+)]	CGC
JT11069	xbx-1(ok279)	CGC, a 1,600-bp deletion
CX3716	lin-15B(n765); kyls141[Posm-9::osm-9::gfp; lin-15(+)]	CGC
GOU1348	casEx1605[Phsp-16.2::Cas9+PU6::che-3 sgRNA; Podr-1::dsRed; unc-76(+)]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1411	casEx5520[Phsp-16.2::Cas9+PU6::lis-1-T1/2 sgRNA; Podr-1::dsRed; unc-76(+)]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1417	casEx5512[Phsp-16.2::Cas9+PU6::dlc-1-T1/2 sgRNA; Podr-1::dsRed; unc-76(+)]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1418	casEx5514[Phsp-16.2::Cas9+PU6::dyci-1-T1/2 sgRNA; Podr-1::dsRed; unc-76(+)]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1419	casEx5513[Phsp-16.2::Cas9+PU6::dl-1-T1/2 sgRNA; Podr-1::dsRed; unc-76(+)]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1421	casEx5516[Phsp-16.2::Cas9+PU6::dylt-3 sgRNA; Podr-1::dsRed; unc-76(+)]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1422	casEx5517[Phsp-16.2::Cas9+PU6::dyrb-1 sgRNA; Podr-1::dsRed; unc-76(+)]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1423	casEx5534[Phsp-16.2::Cas9+PU6::osm-1-T1/2 sgRNA; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1442	casls482[Phsp-16.2::Cas9+PU6::dyci-1 sgRNA T1/2; Podr-1::dsRed (+)]; mnls17	Cross with <i>mnls17</i>
GOU1444	casEx1629[Phsp-16.2::Cas9+PU6::che-11 sgRNA; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1448	casEx1633[Pxbx-1::xbx-1::YFP; rol-6(su1006) (+)]; casls482	Microinjection and cross with <i>casls482</i>
GOU1453	xbx-1(ok279); casEx1630[Pdyf-1::gfp::dyci-1; Pdyf-1::osm-6::mCherry; rol-6(su1006) (+)]	Microinjection
GOU1461	casEx5524[Phsp-16.2::Cas9+PU6::osm-12-T1/2 sgRNA; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24]; mnls17	Microinjection and cross with <i>mnls17</i>
GOU1460	casEx1629[Phsp-16.2::Cas9+PU6::xbx-1 sgRNA; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24]; mnls17	Microinjection
GOU1463	casEx1629[Phsp-16.2::Cas9+PU6::xbx-1 sgRNA; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24]; casls487[Pdyf-1::Cas9; Podr-1::dsRed; unc-76(+)]; mnls17	Cross with <i>casls487</i>
GOU1464	casEx5544[Phsp-16.2::Cas9+PU6::osm-3 sgRNA; Pdyf-1::osm-6::mCherry; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24]	Microinjection
GOU1465	casEx5535[Pdlc-1::dlc-1::GFP; Pdyf-1::osm-6::mCherry; unc-76(+)]	Microinjection
GOU1466	casEx5537[Pdylt-3::dylt-3::GFP; Pdyf-1::osm-6::mCherry; unc-76(+)]	Microinjection
GOU1467	casEx5527[Pdyf-1::GFP::dyci-1; Pdyf-1::osm-6::mCherry; rol-6(su1006) (+)]	Microinjection
GOU1468	casEx5523[Pdyci-1::dyci-1::GFP; Pdyf-1::osm-6::mCherry; rol-6(su1006) (+)]	Microinjection
GOU1469	casEx5538[Pdli-1::dl-1::GFP; Pdyf-1::osm-6::mCherry; unc-76(+)]	Microinjection
GOU1470	casEx5547[Pdyf-1::GFP::dl-1; Pdyf-1::osm-6::mCherry; unc-76(+)]	Microinjection
GOU1471	casEx5548[Pls-1::lis-1::GFP; Pdyf-1::osm-6::mCherry]	Microinjection
GOU1472	casEx5549[Pdyf-1::GFP::lis-1; Pdyf-1::osm-6::mCherry]	Microinjection
GOU1473	casEx5540[Phsp-16.2::Cas9+PU6::osm-3 sgRNA; Pdyf-1::Cas9; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24]; mnls17	Microinjection
GOU1474	casls482[Phsp-16.2::Cas9+PU6::dyci-1 sgRNA T1/2; Podr-1::dsRed (+)]	Microinjection and integration
GOU1475	casls487[Pdyf-1::Cas9; Podr-1::dsRed; unc-76(+)]	Microinjection and integration
GOU1568	casls509; kyls141	Cross with <i>kyls141</i>
GOU1570	casls482; casEx1642[Pdyf-1::gfp::mdyci-1; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24; rol-6(su1006) (+)]	Microinjection
GOU1571	casls509[Phsp-16.2::Cas9+PU6::dlc-1 sgRNA T1/2; Podr-1::dsRed; unc-76(+)]; casEx1643[Pdlc-1::mdlc-1::GFP; Pegl-17::Myri-mCherry; Pegl-17::mCherry::his-24; rol-6(su1006) (+)]	Microinjection and integration
GOU1584	casls482; kyls141	Cross with <i>kyls141</i>

CGC, Caenorhabditis Genetics Center.

Table S2. PCR products for construction of transgenic *C. elegans*

PCR product	5' primer (5' to 3')	3' primer (5' to 3')	Template
Pdylt-3::dylt-3::GFP	CAGCGTGAGTGAGAAGTTGGCTAG	AAGGGCCGTACGGCCGACTAGTAGG	N <sub>2</sub> Genomic DNA

Table S3. Plasmids for construction of transgenic *C. elegans*

Plasmid name	5' primer (5' to 3')	3' primer (5' to 3')	Notes
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dyci-1-T1 sgRNA</i>	GATATTCTCCGTCAAGTGTAGAGCT AGAAATAGC	CTTGACGGAGAATATCTCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dyci-1-T2 sgRNA</i>	TGTACAGACACGGTCAGTTAGAGCT AGAAATAGC	GCACGCTCTGTACACTCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dlc-1-T1 sgRNA</i>	ACCTGGCATTCGATCGTGTAGAGCT AGAAATAGC	CGATGCAATGCCAGTTCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dlc-1-T2 sgRNA</i>	CATCGACTGCCACCCGTTAGAGCT AGAAATAGC	GGTGGCCAGTCGATGCCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dli-1-T1 sgRNA</i>	AGCAAATGCTGTGCTCGTTAGAGCT AGAAATAGC	AAGCACAGCATTGCTCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dli-1-T2 sgRNA</i>	TCGACAGACAGAAGACTGTAGAGCT AGAAATAGC	CTTCTTCGCTGTCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dylt-3 sgRNA</i>	CACAATTCTGTGTCGTTAGAGCT AGAAATAGC	ACACACAGAAATTGTCAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>dyrb-1 sgRNA</i>	CATGATTGCTCCAGACAGTTAGAGCT AGAAATAGC	GTCTGGACCAATCATGATCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>lis-1-T1 sgRNA</i>	ACATGAGTTGTCGGAGTTAGAGCT AGAAATAGC	TCCGACAAACTCATGTTCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>lis-1-T2 sgRNA</i>	GATATCAAGCCACTAGGGTTAGAGCT AGAAATAGC	CTAGTGGCTTGATATCATCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>osm-3 sgRNA</i>	GGCACTGTGTTGCCTAGTTAGAGCT AGAAATAGC	AGGCAAACACAGTGCATCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>xbx-1 sgRNA</i>	CGAGTCATTCGATTGTTAGAGCTA GAAATAGC	ATGCGAAATGACTCGTCAAGACATCTCG AATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>che-3 sgRNA</i>	AGTCGGGATTTCCAATTGTTTAGAGCT AGAAATAGC	TTGGAAATCCGACTCAAGACATCTCG AATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>che-11 sgRNA</i>	GTAGATTGGAGCCACGAGTTAGAGCT AGAAATAGC	GTGGCTCAAATCTACTCCAAGACATCTCG CAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>osm-1-T1 sgRNA</i>	AGTTCCCTCCACAATTGTTAGAGCT AGAAATAGC	TTGTACGGAGGAACCTGCAAGACATCTCG CAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>osm-1-T2 sgRNA</i>	TGCCAGAACAGGTGCCGTTAGAGCTA GAAATAGC	GCACCTCTCTGGCAGACAAGACATCTCG CAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>osm-12-T1 sgRNA</i>	ACTGATTTCGCTCAACTGTTAGAGCT AGAAATAGC	CTTGAGCGAAATCAGTCCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDD162- <b>Pssp-16.2::Cas9+ PU6::</b> <i>osm-12-T2 sgRNA</i>	ATACCCCTGTCATCGTGTAGAGCT AGAAATAGC	CGATTGAACAGGTATCCAAGACATCTC GCAATAGGAGG	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pDONR- <b>Pdfy-1::Cas9</b>	TCTAAGCTTGTCAAATGACAAAAAT ACAGCATCG	GAAGACTAATTGACTTACGCCACTCT GGGACGT	PCR from pDD162- <b>Pssp-16.2::Cas9+ PU6::Empty sgRNA</b>
pPD95.77- <b>Pdyci-1::dyci-1::GFP</b>	GTACCGTAGAAAAACGGTAGATTCTGC AAGTTTCCGTG	TTCTTCTCTTACTCATGTTATAATTCT CCGCTTCTCATTG	PCR from N <sub>2</sub>
pPD95.77- <b>Pdlc-1::dlc-1::GFP</b>	AGTGACCTGTTGCTCCCTTGGAGC CTCAATCGGT	CCCTCCACCTCCGCCCTCACCTCCAGACT TGAATAGCAGGATGGCGA	PCR from N <sub>2</sub>
pPD95.77- <b>Pdli-1::dl-1::GFP</b>	AGTGACCTGTTGCTCAATGGAAACA GTTTGTAGATCGGATTGAC	CCCTCCACCTCCGCCCTCACCTGCATCAC TGTCCCCGGGGTGAGG	PCR from N <sub>2</sub>
pPD95.77- <b>Plis-1::lis-1::GFP</b>	AGTGACCTGTTGCTTGTGACACAA TATTTCACGACC	CCCTCCACCTCCGCCCTCACACGGCATT CCCAAACTTTCAC	PCR from N <sub>2</sub>
pDONR- <b>Pdfy-1::GFP::dyci-1</b>	GATGAACATACAAATGTCAGAACTGA GGAAACTCGAA	GAAGAGTAATTGGACCATGTTATAATTCT CCGCTTCTCA	PCR from N <sub>2</sub>
pDONR- <b>Pdfy-1::dl-1::GFP</b>	TGTAAGCTTGTCAAATGCCACCAACTG CGCAACCACTGG	CCCTCCACCTCCGCCCTCACCTGCATCAC TGTCCCCGGGGTGAGG	PCR from N <sub>2</sub>
pDONR- <b>Pdfy-1::GFP::lis-1</b>	GATGAACATACAAATGAGTTGCGG AGAGGCAAAAG	GAAGAGTAATTGGACTCAACGGCATTCCC AAACTTTG	PCR from N <sub>2</sub>
pDONR- <b>Pdlc-1::dlc-1::GFP</b>	AGTGCACCTGTTGCTCCCTTGGAGC CTCAATCGTA	CCCTCCACCTCCGCCCTCACCTCCAGACT TGAATAGCAGGATGGCGA	PCR from N <sub>2</sub>
pDONR- <b>Pdfy-1::GFP::dyci-1<sup>mT1</sup></b>	ACCTGTCATGTCATGCTTCGACCTCAT TCGATGAAAG	CATATTGAGACAGGTCGGTATTCAACTG AGCCAACAG	PCR from pDONR-Pdfy-1::GFP:: <i>mdyci-1</i>
pDONR- <b>Pdfy-1::GFP::dyci-1<sup>mT1+mT2</sup></b>	GTACAAGCGTTGGACGCTGGAAAATT CCAAGTCGAC	CCAAACCGCTTGTACTGATAACAAAGATT CGCTTAAGC	PCR from pDONR-Pdfy-1::GFP:: <i>dyci-1<sup>mT1</sup></i>
pDONR- <b>Pdlc-1::dlc-1<sup>mT1</sup>::GFP</b>	GTGTTGCACAACTATTGGCTCTGTTG CATGTCATC	TAGATTGTCACACAAGCCCTCGAGAAA TACAAACAT	PCR from pDONR-Pdlc-1::dlc-1:: GFP
pDONR- <b>Pdlc-1::dlc-1<sup>mT1+mT2</sup>::GFP</b>	TATACAGTGCATGTGGGGTTGATTTC TTGTCGAAC	ACATGGCAGTGTAGTAGGAAGAAACTT TGGAACTAC	PCR from pDONR-Pdlc-1::dlc- 1 <sup>mT1</sup> ::GFP

Promoters are highlighted in bold.

Table S4. Targets of CRISPR in *C. elegans*

Gene	CRISPR–Cas9 targets (POM)	
	Target	Sequence
<i>dyci-1</i>	T1	AGGATATTCTCCGTCAAGTAGG
	T2	AGTGTACAGACAGCGTGCACGG
<i>dlc-1</i>	T1	CAACCTGGCATTGCATCGTCGG
	T2	GCCATCGACTGCCACCCAGG
<i>dli-1</i>	T1	ACAGCAAATGCTGTGCTTCGG
	T2	CTTCGACAGACGAAGAAAGTCGG
<i>dylt-3</i>	T1	TACACAATTCTGTGTGTCGG
<i>dyrb-1</i>	T1	ATCATGATTGCTCCAGACAAGG
<i>lis-1</i>	T1	CAACATGAGTTGTCGGAGAGG
	T2	ATGATATCAAGCCACTAGGAGG
<i>osm-3</i>	T1	ATGGCACTGTGTTGCCTATGG
<i>xbx-1</i>	T1	ACGAGTCATTCGCATTTGG
<i>che-3</i>	T1	GAAGTCGGGATTCCAATTGG
<i>osm-1</i>	T1	CAAGTTCCCTCGTACAATTGG
	T2	TCTGCCAGAAGAGGTGCCTGG
<i>che-11</i>	T1	GAGTAGATTGGAGCCACGACGG
<i>dyci-1</i>	mT1	AAGACATATTGAGACAGGT <u>CGG</u>
	mT2	ACCGTCCAACC <u>GGCTTGACTG</u>
<i>dlc-1</i>	mT1	CCACATGGCACTGTATA <u>AGTAGG</u>
	mT2	GCCATCGACTGCCAC <u>ACAAG</u>

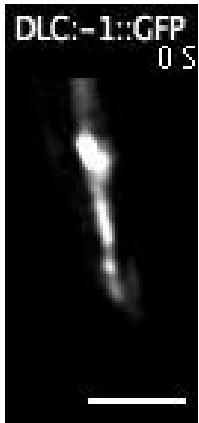
Bold text indicates PAM sequences. Underlined text indicates synonymous mutations of the CRISPR–Cas9 targets for the rescue experiments. Sequences are 5' to 3'.

Table S5. Primers for molecular analysis

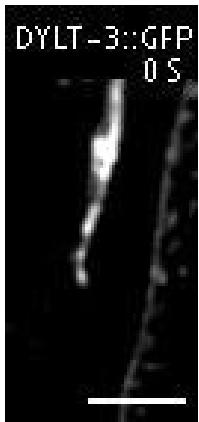
Target gene	5' sequence (5' to 3')	3' sequence (5' to 3')
<i>osm-3</i>	CAAGGACTTCACGTTGATGGAGC	GTCAAAGAGTCAGGTCAAGGGC
<i>xbx-1</i>	CACTATCAGAACGTTCGTGT	CACCAATACAAGTCTAACGCTAG
<i>dyci-1</i>	GGCAGAGCTGAATCTCAGCGG	CGCTCATCGTCAGTCTGGTTCC
<i>dlc-1</i>	GGTTGACCGCAAGGCTGTGATCAA	GCACACTGAAGATCTAACGCCACC
<i>dli-1</i>	GCACGTGGAGCACACGATATCC	GCAGACGAACCTCTCCGAGGG
<i>dylt-3</i>	TGCCATGACGGACCGGAAATACTTT	ACTCATTGCTTTACCGATTCTGCC
<i>dyrb-1</i>	TCGTTCTCGATTCACTGGCTGC	ACGTGGAGGGACACTGTTCTAACGCA
<i>lis-1</i>	CTGCAAAAAATCTGAATTCCCTTTCAATCTC	GATATTTCATCTCTGAACCTTAATGACCAGG



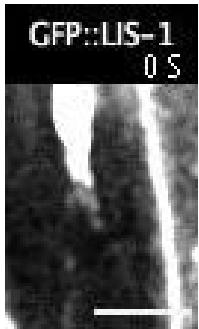
Video 1. IFT of GFP::DYCI-1. Transgenic *C. elegans* strain (GOU1467) expressing GFP-tagged DYCI-1 in cilia. Images were taken using a time-lapse fluorescence microscope (Axio Observer Z1 microscope; Carl Zeiss) attached to a spinning disk confocal scan head (CSU-X1 Spinning Disk Unit; Yokogawa Electric Corporation). Frames were taken continuously for 1.5 min. The display rate was 7 frames per second.



Video 2. **IFT of DLC-1::GFP.** Transgenic *C. elegans* strain (GOU1465) expressing GFP-tagged DLC-1 in cilia. Images were taken using a time-lapse fluorescence microscope (Axio Observer Z1 microscope; Carl Zeiss) attached to a spinning disk confocal scan head (CSU-X1 Spinning Disk Unit; Yokogawa Electric Corporation). Frames were taken continuously for 1.5 min. The display rate was 7 frames per second.



Video 3. **IFT of DYLТ-3::GFP.** Transgenic *C. elegans* strain (GOU1466) expressing GFP-tagged DYLТ-3 in cilia. Images were taken using a time-lapse fluorescence microscope (Axio Observer Z1 microscope; Carl Zeiss) attached to a spinning disk confocal scan head (CSU-X1 Spinning Disk Unit; Yokogawa Electric Corporation). Frames were taken continuously for 1.5 min. The display rate was 7 frames per second.



Video 4. **IFT of GFP::LIS-1.** Transgenic *C. elegans* strain (GOU1472) expressing GFP-tagged LIS-1 in cilia. Images were taken using a time-lapse fluorescence microscope (Axio Observer Z1 microscope; Carl Zeiss) attached to a spinning disk confocal scan head (CSU-X1 Spinning Disk Unit; Yokogawa Electric Corporation). Frames were taken continuously for 1.5 min. The display rate was 7 frames per second.