Functional analysis of an individual IFT protein: IFT46 is required for transport of outer dynein arms into flagella

Yuqing Hou  
*University of Massachusetts Medical School, Yuqing.Hou@umassmed.edu*

Hongmin Qin  
*Yale University*

John A. Follit  
*University of Massachusetts Medical School Worcester, John.Follit@UMassmed.edu*

See next page for additional authors

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Gregory J. Pazour∗∗ Joel L. Rosenbaum†† George B. Witman‡‡

∗University of Massachusetts Medical School, Yuqing.Hou@umassmed.edu
†University of Massachusetts Medical School, John.Follit@UMassmed.edu
∗∗University of Massachusetts Medical School, Gregory.Pazour@umassmed.edu
††University of Massachusetts Medical School, George.Witman@umassmed.edu
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Functional analysis of an individual IFT protein: IFT46 is required for transport of outer dynein arms into flagella

Yuqing Hou,1 Hongmin Qin,3 John A. Follit,2 Gregory J. Pazour,2 Joel L. Rosenbaum,3 and George B. Witman1

1Department of Cell Biology and 2Program in Molecular Medicine, University of Massachusetts Medical School, Worcester, MA 01655
3Department of Molecular, Cellular, and Developmental Biology, Yale University, New Haven, CT 06520

Introduction

Intraflagellar transport (IFT), which is the bidirectional movement of particles within flagella, is required for flagellar assembly. IFT particles are composed of ~16 proteins, which are organized into complexes A and B. We have cloned Chlamydomonas reinhardtii and mouse IFT46, and show that IFT46 is a highly conserved complex B protein in both organisms. A C. reinhardtii insertional mutant null for IFT46 has short, paralyzed flagella lacking dynein arms and with central pair defects. The mutant has greatly reduced levels of most complex B proteins, indicating that IFT46 is necessary for complex B stability. A partial suppressor mutation restores flagellar length to the ift46 mutant. IFT46 is still absent, but levels of the other IFT particle proteins are largely restored, indicating that complex B is stabilized in the suppressed strain. Axonemal ultrastructure is restored, except that the outer arms are still missing, although outer arm subunits are present in the cytoplasm. Thus, IFT46 is specifically required for transporting outer arms into the flagellum.

Correspondence to George Witman: george.witman@umassmed.edu
Abbreviation used in this paper: IFT, intraflagellar transport.

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In *C. reinhardtii*, IFT172 interacts with EB1, which is located at the tip of the flagellum (Pedersen et al., 2003, 2005), and an IFT172 temperature-sensitive mutant is defective in IFT particle turnaround at the tip of the flagellum (Pedersen et al., 2005); based on these results, it was proposed that IFT172 is involved in regulating the transition between anterograde and retrograde IFT (Pedersen et al., 2005). In mammalian cells, IFT20 was found to be unique among IFT particle proteins examined in that it is localized to the Golgi apparatus, as well as the cilium, and moderate knockdown of IFT20 reduced the amount of the membrane protein polycystin-2 in the cilium (Follit et al., 2006), suggesting that IFT20 is involved in trafficking of ciliary membrane proteins. Other than this, nothing is known about the specific roles of the IFT particle proteins and how, or even whether, they interact with specific cargos.

To obtain more information on the function of a specific IFT particle protein, we have focused on IFT46, which was previously briefly reported to be a complex B protein in *C. reinhardtii* (Cole et al., 1998). We have cloned and characterized IFT46 from *C. reinhardtii* and mouse, and find that it is a homologue of DYF-6, a protein very recently reported to undergo IFT in *Caenorhabditis elegans* and to result in truncated dendritic cilia when mutated in the worm (Bell et al., 2006). We also describe the phenotype of a *C. reinhardtii*-null mutant for IFT46, and of a suppressor of that mutant. The observations provide new insights into the role of IFT46 in IFT particle stability and transport of a specific IFT cargo. Portions of this work were previously reported in abstract form (Hou et al., 2005).

### Results

#### Cloning of *C. reinhardtii* IFT46

The gene and cDNA encoding *C. reinhardtii* IFT46 were cloned as described in Materials and methods. The cDNA (accession no. DQ787426) contains a 1,035-nt ORF predicted to encode a 37.9-kD protein (Fig. 1 A) with a pl of 4.61. The cDNA has a stop codon 18 nt upstream of the predicted start codon, and a polyA consensus sequence at nt 1,703–1,707. ESTs have a polyA tail 12–14 nt downstream of the polyA consensus sequence. Therefore, the ORF is complete. No structural domains or motifs were identified within the sequence.

IFT46 was initially identified as an IFT complex B protein, based on its cosedimentation with other complex B proteins on sucrose gradients. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient. The fractions were analyzed by Western blotting using antibodies to IFT46, complex B proteins IFT172 and IFT81, and complex A protein IFT139. Under these experimental conditions, complex B separated from complex A, and IFT172 dissociated from complex B.

### Table I. *C. reinhardtii* IFT particle proteins

<table>
<thead>
<tr>
<th>Complex A</th>
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<tr>
<td>IFT144</td>
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<td>IFT140</td>
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<td>IFT57/55</td>
<td>IFT27</td>
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Figure 1. Identification and characterization of *C. reinhardtii* IFT46.

(A) Predicted amino acid sequence of IFT46 and alignment with orthologues from other organisms. CrIFT46 is highly conserved from aa 100–315. Amino acid identities are marked with asterisks; similarities are marked with either one or two dots. Sequences used in this alignment are as follows: human (AAH22856), mouse (NP_076320), zebrafish (XP_694278), honey bee (XP_396519), and *CrIFT46* (DQ787426). (B) The antibody to IFT46 specifically recognizes a 46-kD doublet in Western blots of whole cell lysates; the doublet is caused by phosphorylation (unpublished data). (C) IFT46 has a typical cellular localization for an IFT particle protein. Cells were labeled with the anti-IFT46 antibody. Images of the same cell were acquired focusing at the flagella (a and c), or at the basal body region (b and d) with much less exposure time. The majority of IFT46 is located at the peribasal body region, with a lesser amount distributed along the flagella as distinct dots. Bar, 5 μm. (D) IFT46 comigrates with IFT81, an IFT complex B protein, in sucrose gradients. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient. The flagellar membrane plus matrix was fractionated in a 5–20% sucrose gradient.

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in sucrose gradients (Cole et al., 1998). Characterization of our cloned protein indicates that it behaves exactly as expected of a 46-kD complex B protein, as follows: a) six unique peptides corresponding to the cloned protein were identified in the membrane plus matrix fraction of the flagellar proteome, but no peptides from it were found in any other fraction, which is a distribution typical for IFT particle proteins but unusual for non-IFT proteins (Pazour et al., 2005); b) using real-time PCR, we found that expression of the protein is up-regulated 10.4 ± 2.03-fold (SEM) upon deflagellation, which is characteristic of flagellar proteins, including other complex B proteins (Pazour et al., 2005); c) an antibody to a peptide contained in the cloned protein was generated and shown to react specifically with a single protein of M₉, ~46,000 in Western blots of whole cell lysates (Fig. 1 B); d) immunofluorescence microscopy with the antibody as probe showed that the majority of the cloned protein is located in the basal body region, with a lesser amount in puncta along the length of the flagella (Fig. 1 C), is a distribution identical to that of other IFT particle proteins (Cole et al., 1998; Deane et al., 2001); moreover, the protein colocalized with complex B protein IFT172, but not complex A protein IFT139 (see the section Complex A and B proteins are located in distinct compartments in the basal body region; Fig. 6 B); and e) when the flagellar membrane plus matrix fraction was analyzed by sucrose density gradient centrifugation, the cloned protein is located in the basal body region, with a lesser amount in the cilia. The inset shows an enlargement of one of the cilia. Bar, 10 μm.

IFT complex B proteins coimmunoprecipitate with MmIFT46. Whole-cell lysates from mouse IMCD3 cells expressing Flag-tagged MmIFT46 were labeled with antibodies to Flag (red), GFP (green), which localizes to Golgi in addition to cilia, and a centrosome component (blue). Nuclei were DAPI stained and are shown as gray. IFT46-Flag was concentrated at the bases of the cilia, but was also located along the length of the cilia. The inset shows an enlargement of one of the cilia. Bar, 10 μm.

IFT46 is required for transport of outer arm dynein

To investigate the role of IFT46 in intraflagellar transport, we screened a collection of CrifTF46 insertional mutants by Southern blotting and identified one strain, T8a44-11, with a defect in the ift46 gene. Analysis by PCR showed that the mutant allele, which we term ift46-1, has a deletion or disruption somewhere between the fourth and the seventh exon of the ift46 gene (Fig. 3, A and B). This mutant has short, stumpy flagella.

Strain T8a44-11 was backcrossed to wild-type cells, and a progeny, YH6, which lacked the pJ mutation carried by the parental strain, was selected for detailed characterization. As in T8a44-11, the IFT46 gene in YH6 is disrupted, as shown by Southern blotting (Fig. 3 C, lane 2). No IFT46 can be detected in lysates of YH6 cells (Fig. 3 D, lane 2), indicating that IFT46 is not expressed in YH6 cells. Thus, the mutant allele is a null allele (see also the section The 3′ end of the IFT46 gene is
transcribed in Sup13461 cells). Like T8a4-11, YH6 cells have short, stumpy flagella that barely extend beyond the flagellar collar (Fig. 4). The flagella are nonmotile.

To confirm that the mutant phenotype of YH6 was caused by the disruption of the IFT46 gene, YH6 cells were transformed with a 4.8-kb fragment containing only the wild-type IFT46 gene (Fig. 3 A). Numerous wild-type swimmers with full-length flagella were recovered. Southern blotting revealed that the exogenous IFT46 gene had integrated into the genome at different sites in several of these rescued strains (Fig. 3 C, lanes 3–9), confirming that the rescued strains were independently derived. Therefore, restoration of motility was caused by incorporation of the transgene, and not caused by disruption of some other gene. Western blotting confirmed that expression of IFT46 was restored in the transformants (Fig. 3 D, lanes 3–9). These results demonstrate that the short flagella phenotype of YH6 is caused by the absence of IFT46. Hereafter, this strain will be referred to as the ift46 mutant.

The ift46 mutant has unique defects in its axoneme

Although very short, ift46 flagella are longer than those of mutants with defects in the complex B proteins IFT88 (Pazour et al., 2000) and IFT52 (Brazelton et al., 2001), which do not form flagella beyond the transition region. Because flagella are formed in the ift46 mutant, we were able to compare them with flagella from wild-type and rescued cells by EM to identify flagellar defects associated with loss of IFT46. Serial sections of wild-type flagella have shown that the outer doublet microtubules are connected by rodlike “peripheral links” in the most proximal part of the flagella (Fig. 4 A, a); the rows of dynein arms begin at a level slightly more distal, but still within the flagellar collar (Fig. 4 A, b; Hoops and Witman, 1983). The ift46 flagella have nine outer doublet microtubules and frequently extend to the limits of the flagellar collar or beyond (Fig. 4 A, f–p), but we never observed dynein arms in longitudinal or cross sections of these axonemes. In addition, the mutant flagella lack the projections into the lumens of the B-tubule (Hoops and Witman, 1983) and frequently have defects in the central pair of microtubules (Fig. 4 A, f–k). In contrast to mutants with defects in the retrograde IFT motor (Pazour et al., 1999; Porter et al., 1999; Hou et al., 2004), few, if any, IFT particles accumulate in the ift46 mutant flagella. It is important to note that the flagella from the rescued cells (Fig. 4 A, d and e) have typical wild-type morphology with normal inner and outer dynein arms and central pair microtubules; this confirms that
IFT46 is required for transport of outer arm dynein

The ultrastructural defects seen in the mutant are caused by loss of IFT46.

To determine whether the dynein arm deficiency in the ift46 mutant is caused by degradation of dyneins within the cell body, by an inability to transport dyneins into the flagella, or by an inability to assemble them onto the axonemes, we analyzed whole cell lysates and flagella from ift46 and wild-type cells by Western blotting (Fig. 4 B). The cell lysates of the mutant contained normal levels of the outer arm dynein intermediate chain IC2 and the inner arm dynein intermediate chain IC138. Therefore, the dyneins are present in the mutant cells. However, both IC2 and IC138 were completely absent from the ift46 mutant flagella, indicating that the dyneins are not transported into the flagella. In contrast, DC2, which is a component of the outer dynein arm docking complex that is transported into the wild-type flagellum and assembled onto the doublets independently of the outer dynein arm (Wakabayashi et al., 2001), is transported into the ift46 flagella. The presence of DC2 in the flagella provides additional evidence that the lack of dynein arms is not simply because of the short length of the mutant flagella or to a general failure to transport proteins into the flagellum. These data confirm that the ultrastructural findings that the ift46 mutant has a defect in transporting dynein arms into the flagella.

IFT complex B is unstable in the absence of IFT46

To investigate the role of IFT46 in IFT complex assembly, we examined the cellular levels of IFT complex B and A proteins in cell lysates of wild-type and ift46 cells (Fig. 5, lanes WT and ift46). When normalized with tubulin, the levels of complex B proteins IFT20, IFT57, IFT72, and IFT81 were greatly decreased in the mutant cells relative to the wild-type cells. The only complex B protein not reduced in the absence of IFT46 was IFT172, the level of which was the same as or greater than in wild-type
Suprfl461 cells, and mutant were increased by protein levels are increased in antibodies to several IFT particle proteins, IC2, and DC2. Complex A proteins IFT172 or complex A into or out of the flagellum, transport of IFT172 or complex A proteins in wild-type and PCR to measure transcript levels for several complex A and B proteins in wild-type and mutant (Fig. 6 C). These results support the hypothesis that a small number of incomplete complex B particles assemble from the residual complex B proteins and are capable of being transported into the flagellum in the absence of IFT46. The resulting low level of IFT may account for the ability of the jfl46 cells to assemble their short flagella.

Loss of IFT46 is specifically correlated with loss of the outer dynein arm in a partially suppressed strain

jfl46 cells are between those in wild-type and Suprfl461 cells have IC2 and DC2. Complex A and B proteins are located in distinct compartments in the basal body region

To determine if the absence of IFT46 and the accompanying large decrease in most other complex B proteins affected the transport of IFT172 or complex A into or out of the flagellum, we used immunofluorescence microscopy to examine jfl46 cells that were double labeled with antibodies to tubulin and IFT172 or IFT139 (Fig. 6 A). In both cases, the IFT particle proteins were concentrated around the basal bodies and in the short flagella. Thus, both proteins are transported into the flagella in the absence of IFT46. Surprisingly, however, the distributions of IFT172 and IFT139 differed from each other in the cell body, with that of IFT172 (Fig. 6 A, a–d) appearing to have almost no overlap with that of IFT139 (Fig. 6 A, e–h), which was more anterior and often concentrated into two distinct lobes.

To clarify whether this difference in distribution was normal or caused by loss of IFT46, wild-type cells were double labeled with antibodies to IFT46 and IFT172 or IFT139. In most cases, IFT46 and IFT172 colocalized precisely with each other in the peribasal body region (Fig. 6 B, a–c), which is consistent with the other evidence that IFT46 is a complex B protein. In contrast, although IFT139 colocalized with IFT46 at the extreme apical end of the cell, the labeling of IFT46 almost always extended more posteriorly than that of IFT139 (Fig. 6 B, d–f). This is the first observation that complex A and B proteins differ in their distribution, and indicates that the complexes, or at least a subset of them, are not physically associated in the cell body.

Our observation that IFT172 was transported into the short flagella of the jfl46 mutant (Fig. 6 A) raised the question of whether IFT172 was being transported into the flagella independently of other complex B proteins or in association with incomplete complex B particles, possibly assembled from the small amount of complex B proteins still present in the mutant. To address this question, we used immunofluorescence microscopy to examine the distribution of another complex B protein, IFT57, the levels of which are greatly reduced in the mutant. Like IFT172, the residual IFT57 was transported into the short flagella of the jfl46 mutant (Fig. 6 C). These results support the hypothesis that a small number of incomplete complex B particles assemble from the residual complex B proteins and are capable of being transported into the flagellum in the absence of IFT46. The resulting low level of IFT may account for the ability of the jfl46 cells to assemble their short flagella.

cells, depending on the preparation, suggesting that the level of IFT172 is controlled independently from that of the other complex B proteins. In contrast to the decrease in most complex B proteins seen in the jfl46 mutant, the levels of complex A proteins IFT139 and IFT140 were greatly increased in jfl46 cells relative to wild-type cells.

To examine if these differences in protein levels were caused by changes in synthesis or stability, we used real-time PCR to measure transcript levels for several complex A and B proteins in wild-type and jfl46 cells. Transcript levels in the mutant were increased by ~2.8- and ~2.0-fold for complex A proteins IFT140 and IFT139, and by ~1.8- and ~1.6-fold for complex B proteins IFT81 and IFT72, respectively. Therefore, the mutant responds to its defect by increasing the mRNA levels of at least these complex A and B proteins. The increase in complex A proteins in the mutant presumably reflects this increase in mRNA abundance. However, because the levels of most complex B proteins are drastically decreased in the mutant, even though complex B mRNA levels in general appear to be increased, it is likely that these proteins are specifically degraded in the absence of IFT46.

Figure 5. IFT complex B is unstable in the jfl46 mutant, but stabilized in the partially suppressed strain. Whole-cell lysates from wild-type cells, Suprfl461 cells, and jfl46 cells were analyzed in Western blots probed with antibodies to several IFT particle proteins, IC2, and DC2. Complex A protein levels are increased in jfl46 compared with wild-type cells. With the exception of IFT172, complex B protein levels are dramatically decreased in jfl46 compared with wild-type cells. The levels of these proteins in Suprfl461 cells are between those in jfl46 and wild-type cells. IFT172 is controlled independently from that of the other complex B proteins. In contrast to the decrease in most complex B proteins IFT81 and IFT72, respectively. Therefore, the mutant responds to its defect by increasing the mRNA levels of at least these complex A and B proteins. The increase in complex A proteins at the extreme apical end of the cell is lost. This, together with the results for the jfl46 mutant, also shows that in the absence of IFT46, the colocalization of IFT172 with the other evidence that IFT46 is a complex B protein. In contrast, although IFT139 colocalized with IFT46 at the extreme apical end of the cell, the labeling of IFT46 almost always extended more posteriorly than that of IFT139 (Fig. 6 B, d–f). This is the first observation that complex A and B proteins differ in their distribution, and indicates that the complexes, or at least a subset of them, are not physically associated in the cell body.
proteins in stimulated Supif461, wild-type, and ift46 cells were compared by Western blotting. In the Supif461 cells, complex B proteins IFT20, IFT57, IFT72, and IFT81 were increased to a level between those of ift46 and wild-type cells (Fig. 5), whereas the levels of the complex A proteins IFT139 and IFT140 were decreased to a level between those of ift46 and wild type. Importantly, IFT46 is still undetected in the suppressed strain. This result indicates that the partial suppression of ift46 involves an increased stability of IFT complex B in the absence of full-length IFT46. It is possible that a C-terminal fragment of IFT46 is expressed in Sup ift461 cells and incorporated into complex B, thereby stabilizing it. Such a fragment would not be detected by our antibody to the N terminus of IFT46.

The slow, jerky swimming of Supif461 is typical of outer dynein arm mutants. Therefore, we used immunofluorescence microscopy to check for the presence of outer and inner dynein arms in Supif461 flagella. No outer arm dynein was detected using an antibody to the α heavy chain of outer arm dynein (Fig. 7 A, a–i). In contrast, labeling of Supif461 flagella by an antibody to inner arm dynein I1 intermediate chain IC138 was normal (Fig. 7 A, j–r). These results show that transport into the flagellum of inner arm dynein I1, but not outer arm dynein, is restored in the suppressed strain.

To determine the extent to which the ultrastructural defects of ift46 were restored in the partially suppressed strain, Supif461 cells and flagella were examined by electron microscopy (Fig. 7 B). The inner arms, radial spokes, and central microtubules were present and appeared normal. However, no outer dynein arms were observed. Therefore, the suppressor strain assembles flagella that lack the outer dynein arm but appear normal in every other way. Western blotting showed that the levels of both outer arm dynein and outer dynein arm docking complex proteins, as

Figure 6. Complex A and B proteins differ in cellular distribution, and loss of IFT46 affects the cell body localization of IFT172, but not of IFT139. (A) ift46 cells were double labeled either with antibodies to IFT172 (a) and tubulin (b) or with antibodies to IFT139 (e) and tubulin (f). c and g are merged images; d and h are enlargements of the flagella and basal body regions. IFT172 and IFT139 are both present in the short flagella (arrows). In the basal body region, IFT139 localizes more anteriorly than IFT172. (B) Wild-type cells were double-labeled either with antibodies to IFT172 (a) and IFT46 (b) or with antibodies to IFT139 (d) and IFT46 (e). The merged images are shown in c and f; the insets show enlargements of the basal body regions. IFT172 usually colocalizes with IFT46. However, IFT139 usually only partially colocalizes with IFT46 in the anterior part of the basal body region. (C) In the absence of IFT46, the remaining complex B proteins are still transported into the flagella. ift46 cells were labeled with an antibody to IFT57, which is located in flagella (arrows), as well as in the basal body region. Bars, 5 μm.
represented by IC2 and DC2, respectively, were normal in Supift461 cells (Fig. 5). Thus, the inability to transport and assemble outer arms in the flagella is not simply caused by an absence of these components from the cell cytoplasm. These results indicate that IFT46 is specifically needed to transport outer dynein arm components into the flagellum. The inner dynein arm and central pair defects observed in the ift46 mutant are likely attributable to a more general deficiency in IFT caused by the reduced number of complex B particles.

The 3′ end of the IFT46 gene is transcribed in Supift461 cells

Analysis by PCR revealed that the suppressor mutation involved a rearrangement or deletion somewhere in the region between the 3′ end of the inserted NIT1 gene and the seventh intron of the IFT46 gene (unpublished data). To determine if this mutation caused a change in the transcription of the IFT46 gene, wild-type, ift46, and Supift461 cells were examined by real-time PCR using primer pairs designed to assay for the presence of the 5′ end, middle, and 3′ end of the IFT46 mRNA (Fig. 8). In wild-type cells, all three regions were detected. In the ift46 mutant, only the 5′ end was detected, indicating that the 5′ end of the gene is transcribed, but a full-length mRNA is not made. Because our antibody to the N terminus of IFT46 did not detect a product, it appears that the truncated mRNA is not translated into a stable protein. This provides further evidence that ift46 is a null allele. In Supift461 cells, both the 5′ and the 3′ end, but not the middle, were reproducibly detected. Therefore, the suppressor mutation results in transcription, and possibly translation, of the 3′ end of the IFT46 gene. Our antibody would not detect a product containing the C-terminal end of IFT46, but lacking its N-terminal end. However, we can rule out the possibility that the suppressor mutation results in translation of the N-terminal part of IFT46 fused with the C-terminal part of IFT46, because our antibody did not detect any product in Supift461 cells. Transcripts encoding the 3′ end of the IFT46 gene were detected in Supift461 cells in both the presence and absence of aeration, so the suppressor mutation, not stress, causes transcription of the 3′ end of the gene.

Discussion

The C. reinhardtii IFT46 sequence, which is reported for the first time in this study, demonstrates that this protein, like other IFT particle proteins, is highly conserved among ciliated organisms. C. reinhardtii IFT46 was previously reported to cosediment with other IFT complex B proteins (Cole et al., 1998). We confirm that IFT46, in both C. reinhardtii and mammals, is a complex B protein based on cosedimentation, coimmunolocalization, and coimmunoprecipitation with other complex B, but not complex A, proteins. Our analysis of a C. reinhardtii ift46 mutant and a suppressed strain of the mutant indicate that IFT46 is necessary for complex B stability and is specifically required to transport outer dynein arm complexes into the flagella.

Our sequencing of C. reinhardtii IFT46 also revealed that it is a homologue of C. elegans DYF-6 (Starich et al., 1995), the sequence of which was recently reported by Bell et al. (2006).
IFT46 is required for transport of outer arm dynein into flagella

In addition to being very short, ift46 flagella have defects in central pair and dynein arm assembly. Aspects of this phenotype are undoubtedly caused by the greatly reduced amount of complex B in the ift46 mutant. Because there is not enough IFT machinery to transport a full complement of axonemal proteins, the flagella are short and assembly of specific axonemal structures is affected.

However, our discovery of a suppressor mutation that stabilizes complex B in the absence of full-length IFT46 and
restores flagellar assembly, including assembly of the central pair and the inner dynein arms, but does not restore outer dynein arm assembly, even though outer arm proteins are present in the cytoplasm, strongly argues that IFT46 has a specific role in outer dynein arm transport. This is the first direct evidence that the outer dynein arms require IFT for transport into the flagellum, and the first evidence connecting a specific IFT particle protein with a specific cargo. Because the outer arm components are preassembled in the cytoplasm into complexes as large as 1.2 MD (Fowkes and Mitchell, 1998), it is likely that IFT is needed to move them efficiently into the flagellum and out to the flagellar tip, which is the site of axonemal assembly (Witman, 1975; Johnson and Rosenbaum, 1992).

Further work will be necessary to determine if IFT46 is involved directly in outer arm binding, or if loss of IFT46 causes a conformational change in the IFT particle that eliminates an outer arm binding site at some distance from IFT46. However, the fact that the suppressor mutant has a partially stabilized complex B but still fails to transport outer arms into the flagella strongly supports the first possibility. The first possibility is further supported by the recent finding that the human homologue of ODA16, which is a C. reinhardtii flagellar protein that is not a component of dynein but is essential for outer arm transport into the flagellum (Ahmed and Mitchell, 2005), interacts with the mouse homologue of IFT46 in a yeast two-hybrid system, and thus may be an adaptor coupling IFT46 to the outer arm (Ahmed et al., 2006).

Our finding that IFT is involved in transport of outer arm complexes explains a previous observation that antibodies to complex B proteins IFT52 and IFT72 coimmunoprecipitated subunits of the outer dynein arm from a flagellar membrane plus matrix fraction (Qin et al., 2004). An earlier study had reported that inner dynein arms, but not outer dynein arms, required the activity of FLA10, which is one motor subunit of the anterograde IFT motor kinesin-2, for transport into the flagellum (Piperno et al., 1996). There are at least two possible explanations for the difference between these results and our own. First, as discussed by Qin et al. (2004), the earlier studies used a fla10 temperature-sensitive mutant and observations 45–75 min after shift to the restrictive temperature, which may not have been adequate for complete cessation of FLA10 activity. Second and more interestingly, it is possible that there is more than one kinesin for anterograde IFT in C. reinhardtii, as there is in C. elegans (Snow et al., 2004). Phylogenetic analysis on complete kinesin repertoires of a diversity of organisms revealed that some kinesin families are specific for ciliated species (Wickstead and Gull, 2006). The C. reinhardtii kinesin-2 motor subunits FLA10 and FLA8 and the central pair kinesin KLP1, as well as several novel kinesins, were grouped in these subfamilies; two of the latter (C_250150 in the kinesin-9 family, and C_710026 in the newly proposed kinesin-17 family) were each identified by multiple hits to a single peptide (shared with FLA8) in the C. reinhardtii flagellar proteome (Pazour et al., 2005; http://labs.umassmed.edu/chlamyfp/protector_login.php). It is possible that one of the novel kinesins is an IFT anterograde motor, and that it and kinesin-2 transport IFT particles linked to different cargos.

We also observed that the outer dynein arm docking complex, as represented by its DC2 subunit, was transported into the ift46 mutant flagella. The movement of the docking complex, but not the outer dynein arm, into ift46 flagella is consistent with previous results that the docking complex is preassembled in the cytoplasm as a distinct complex not associated with the outer arm (Wakabayashi et al., 2001), and that it is transported into the flagellum independently of the outer arm (Takada and Kamiya, 1994; Wirschell et al., 2004). DC2 was communoprecipitated by antibodies to IFT particle proteins (Qin et al., 2004), indicating that it interacts with the IFT machinery and probably is dependent on it for transport into the flagellum. Our results clearly show that the docking complex does not specifically require IFT46 for entry into the flagellum.

Complex A and complex B occur in distinct, but overlapping, compartments in the basal body region

Complexes A and B are associated with each other in large linear assemblages during transport within the flagellum (Qin et al., 2004), and it has been proposed that turnover of IFT particle proteins and motors at the tip of the flagellum involves the dissociation of complex A from complex B, followed by their reassociation before retrograde transport (Pedersen et al., 2006). However, virtually nothing was known about the interactions of the complexes at the base of the flagellum. Our immunofluorescence microscopy observations have revealed different patterns of localization for complex A and complex B proteins in the wild-type cell, with the former localized more apically in the peribasal body region. In contrast, the complex B proteins IFT172 and IFT46 usually colocalized precisely with each other. These results suggest that complex A and complex B separate from each other upon passage from the flagellum into the cytoplasm, are sorted into separate albeit overlapping compartments, and are subsequently reassembled before their transport into the flagellum, in a reversal of the process proposed to occur at the tip of the flagellum. The region of overlap at the apical end of the peribasal body region may correspond to the basal body transition fibers, which are proposed to be docking sites where the IFT particles are assembled or disassembled before entry into the flagellum or cytoplasm, respectively (Deane et al., 2001; Rosenbaum and Witman, 2002). Within the cell body of the ift46 mutant, the IFT172 in excess over other complex B proteins localizes primarily to the posterior peribasal body region and has little or no overlap with IFT139 in the anterior peribasal body region, indicating that IFT172 does not interact directly with complex A in the cell body.

Our immunofluorescence microscopy studies also showed that both complex A and residual complex B proteins are transported into the short flagella of the ift46 mutant, indicating that even in the absence of suppression, transport of the IFT complexes into the flagella does not require ift46.

Stability of complex B in Supift46 cells may be caused by expression of the C-terminus of IFT46

Our analysis of the Supift46 cells revealed that they differ from ift46 cells in that the suppressor mutation causes transcription of the 3’ end of the IFT46 gene. This could come about as a
result of an intragenic mutation that allows promotion of the 3' end of the IFT46 gene from the NIT1 promoter present in the vector originally used to generate the ift46 insertion mutant. Irrespective of the mechanism, the results suggest that the C-terminal portion of IFT46 is expressed, possibly as part of a fusion protein with nitrate reductase (encoded by the NIT1 gene). This fragment may then be incorporated into complex B, stabilizing it.

Expression of the suppressed phenotype was observed only in Supmt1 cells grown in the absence of aeration. A possible explanation for this is that under stress conditions (hypoxia), a chaperone is produced that helps stabilize complex B. ift46 cells do not form flagella in the absence of aeration, so this hypothetical chaperone does not stabilize complex B in the complete absence of IFT46. It may be that a stress-induced chaperone can stabilize complex B in the presence of a C-terminal fragment of IFT46, but not in the absence of the fragment.

Flagella are formed in Supmt1 cells, but the flagella lack the outer dynein arms. Therefore, if complex B is indeed stabilized by a C-terminal fragment of IFT46 in Supmt1 cells, this would imply that the N-terminal end of IFT46 is essential for transport of outer arms into the flagellum.

Cells can compensate for defects in IFT complex B
Our identification of a partial suppressor of the ift46 phenotype is the second report of suppression of the phenotype resulting from disruption of a complex B protein. Brown et al. (2003) identified a spontaneous partial suppressor of a null mutant of IFT52 in Tetrahymena thermophila. The T. thermophila ift52-null mutant has basal bodies that fail to form flagella or form short flagella that lack the central pair of microtubules, whereas a variable number of cells of the partially suppressed strain had slightly longer flagella, of which ~13% had a central pair of microtubules, depending on growth conditions. Just as the suppressed phenotype of C. reinhardtii Supmt1 cells was stimulated by lack of aeration, the suppressed phenotype of the T. thermophila strain was stimulated by pericellular hypoxia; suppression of the T. thermophila strain was also stimulated by growth at abnormally low temperature. This suppression differed from that which we observed, in that it arose spontaneously with high frequency, whereas ours was a rare event that was observed only once in 2 yr of culturing the cells; however, in both cases, the phenotype was stable during vegetative growth. Therefore, the mechanism of suppression may be similar in both C. reinhardtii and T. thermophila. In our case, Western blotting showed that the IFT protein levels in the partially suppressed strain were restored to levels between those of wild-type cells and the ift46 mutant cells, indicating that restoration of the ability to form flagella was caused by stabilization of complex B in the absence of full-length IFT46.

Materials and methods
Cells and culture media
C. reinhardtii strains 137C (nit1, nit2, mt+), CC124 (nit1, nit2, mt–), and S12D were obtained from the Chlamydomonas Genetics Center (Duke University, Durham, NC). T8a4-11 (ift46::NIT1, nit1, p1, mt+) was generated by K. Kozminski and J. Rosenbaum (Yale University, New Haven, CT) by transforming KK30A3 (nit1, NIT2, p1, mt+) with the plasmid PM124 linearized with EcoRI. YH6 (ift46::NIT1, PF1, mt+) is an offspring of a cross between T8a4-11 and CC124. Supmt1 is a spontaneous partial suppressor for YH6. Cells were grown in M (Sager and Granick [1953] medium I altered to have 0.0022 M KH2PO4 and 0.00171 M K2HPO4), M-N (M medium without nitrogen), or TAP (Gorman and Levine, 1965) media. Murine IMCD3 cells (COSTECH Laboratories, Inc.) were grown as described in Follit et al. (2006).

Antibodies
The antibodies used are listed in Table S1 (available at http://www.jcb.org/cgi/content/full/jcb.200608041/DC1). The rabbit antibody to IFT46 was generated against a synthetic peptide corresponding to the protein's N-terminal 19 amino acids (Pocano Rabbit Farm) and affinity purified using the same 19 amino-acid peptide.

C. reinhardtii IFT46 gene cloning
165 IFT particles were purified from C. reinhardtii flagella (Cole et al., 1998) and the particle proteins separated by PAGE. A band corresponding to IFT46 was excised and microsequenced. Two peptides, VPRDTPKPDYGKK and KPFDYPDAVCGIDFETK, were obtained; these identified the protein C_21300077 predicted by the C. reinhardtii genome (v. 2; http://genome.jgi-psf.org/crg1-bin/search?mrdb+chre2). Several EST clones in Genbank that contain these two peptides in their ORF were used to clone the IFT46 cDNA. A 4.8-kb fragment that contains only the full-length IFT46 gene was cloned from C. reinhardtii genomic DNA after its amplification by PCR with primers IFT46-5 and IFT46-6 (the sequences of all primers are given in Table S2, available at http://www.jcb.org/cgi/content/full/jcb.200608041/DC1) using ELongase Enzyme Mix (Invitrogen). The sequences of the 5'– and 3'-UTRs of the IFT46 cDNA were verified by sequencing the cloned IFT46 gene; the sequence of the coding region was verified by sequencing a PCR product from a cDNA library.

IFT46 homologues were identified by searching the translated nr database at http://www.ncbi.nlm.nih.gov/BLAST/. Sequences were analyzed as described in Hou et al. (2004).

DNA and RNA isolation and analysis
DNA was isolated from C. reinhardtii as described in Pazzou et al. (1998). DNA gel electrophoresis was carried out by standard procedures (Sambrook et al., 1989). Southern blotting was performed using the DIG High Prime DNA Labeling and Detection Starter Kit II (Roche); instead of using the kit's hybridization buffer, we used Church buffer (7% SDS, 1 mM Na2HPO4, pH 7.2; Church and Gilbert, 1984) and hybridized it at 65°C.

IFT46 gene induction upon deflagellation was analyzed by real-time PCR, as described in Pazzou et al. (2005), using primers IFT46-3 and 4. The ratio of the amount of IFT46 message after deflagellation to that before deflagellation was calculated for each trial. Three independent sets of mRNA were isolated and analyzed three times each.

To measure the mRNA levels of IFT140, IFT81, and IFT72, cDNAs were prepared from cells at the mid-log phase of growth, and quantitated by real-time PCR as described in Pazzou et al. (2005) using the primer pairs IFT140F/IFT140R, IFT81F/IFT81R, or IFT72F/IFT72R. Two independent sets of cDNAs were isolated and analyzed three times each.

To assay transcription of the IFT46 gene, cDNAs were prepared from wild-type cells and from ift46 and Supmt1 cells with or without aeration. Real-time PCR was performed using primer pairs to the 5' end of the gene (IFT46-11/IFT46-2), middle part of the gene (IFT46-9/IFT46-28), and 3' end of the gene (IFT46-10/IFT46-11). Samples were normalized using G protein β subunit (Pazzou et al., 2005). The end products were examined on a 1.5% agarose gel. Three independent sets of mRNA were isolated and analyzed three times each.

A collection of insertion mutants having motility defects was screened for a defect in IFT46 by Southern blotting, using an 864-bp partial cDNA fragment amplified by PCR with primers IFT46-G1 and IFT46-G2 as a probe. The mutated region in the IFT46 gene in the mutant was located by PCR using primer pairs 1 (IFT46-1/IFT46-2), 2 (IFT46-9/IFT46-10), and 3 (IFT46-3/IFT46-4).

Murine IFT46 and Flag-tagged proteins
The open reading frame of MmIFT46 was PCR amplified from mouse testis cDNA using primers mIFT46-1 and -2. The PCR product was digested with BamHI, cloned into the BglII site of pGAL131, and called pJAF161. pJAF113.1 was derived from p3XFLAG-myc-CMV26 (Sigma-Aldrich) by

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filling in the HindIII site to shift the polylinker by four nucleotides. pJAF161.24 encodes a fusion protein in which the 3× Flag tag is fused to the N-terminal end of IFT46. pJAF161.41 encoding GFPFlag was constructed by moving the XbaI–EcoRI GFP-containing fragment from pEGFP N2 (CLONTECH Laboratories, Inc.) into pJAF113.1. pJAF134.3 encoding MinIF20Flag was described by Follit et al. (2006).

Immunoprecipitations
IMCD3 cells that had been transfected with JAF161.24 (MinIF46-Flag), pJAF134.3 (MinIF20-Flag), or pJAF161.41 (GFPFlag) were lysed in Cell lytic M (Sigma-Aldrich) containing 0.1% Tween 20 and 0.1% CHAPS (Bio-Rad Laboratories). After incubation for 10 min at 4°C, the extract was clarified by centrifugation at 18,000 g and treated for 10 min with Sepharose-4B beads, which were removed by centrifugation through a Macro Spin Column (Harvard Apparatus). The treated extract was then incubated with anti-Flag M2-Agarose Affinity Gel (Sigma-Aldrich) for 1 h at 4°C. Unbound proteins were removed by washing the beads in Wash Buffer (Sigma-Aldrich) containing 1% Tween 20 and 150 mM NaCl, followed by washes in Wash Buffer alone. Bound proteins were eluted with 200 µg/µl 3× Flag Peptide (Sigma-Aldrich) and analyzed by Western blotting, as described in Pazzour et al. (1998).

Protein biochemistry
Preparation of C. reinhardtii whole-cell extracts, isolated flagella, and the flagellar membrane plus matrix fraction, as well as PAGE and Western blotting, were performed as described in Pazzour et al. (1999). Sucrose gradient analysis was carried out as described by Hou et al. (2004).

Genetic analysis and transformation
Gradient analysis was carried out as described by Hou et al. (2004). IS3000 genomic Southern analysis and in vivo transformation were performed as described in Pazour et al. (1999). Sucrose gradient analysis was carried out as described by Hou et al. (2004). T8a4-11 and CC124 gametes were induced to mate by dibutyryl-cAMP (Pazour et al., 1999; Hou et al., 2004).

Preparation of whole-cell extracts
Whole-cell extracts from C. reinhardtii cells were fixed and stained for immunofluorescence microscopy by Follit et al. (2006). Images were prepared for final publication using Photoshop 6.0 (Adobe).

Online supplemental material
Table S1 shows the antibodies used in this work. Table S2 shows the primer sequences used in this study. The online version of this article is available at http://www.jcb.org/cgi/content/full/jcb.200608041/DC1.

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