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Cycling of the signaling protein phospholipase D through cilia requires the BBSome only for the export phase

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The BBSome is a complex of seven proteins, including BBS4, that is cycled through cilia by intraflagellar transport (IFT). Previous work has shown that the membrane-associated signaling protein phospholipase D (PLD) accumulates abnormally in cilia of Chlamydomonas reinhardtii bbs mutants. Here we show that PLD is a component of wild-type cilia but is enriched ~150-fold in bbs4 cilia; this accumulation occurs progressively over time and results in altered ciliary lipid composition. When wild-type BBSomes were introduced into bbs cells, PLD was rapidly removed from the mutant cilia, indicating the presence of an efficient BBSome-dependent mechanism for exporting ciliary PLD. This export requires retrograde IFT. Importantly, entry of PLD into cilia is BBSome and IFT independent. Therefore, the BBSome is required only for the export phase of a process that continuously cycles PLD through cilia. Another protein, carbonic anhydrase 6, is initially imported normally into bbs4 cilia but lost with time, suggesting that its loss is a secondary effect of BBSome deficiency.

Introduction

Bardet-Biedl syndrome (BBS; OMIM accession no. 209900) is a rare inherited disorder characterized by retinal degeneration, anosmia, kidney anomalies, polydactyly, hypogonadism, and obesity (Beales, 2005; Blacque and Leroux, 2006; Zaghloul and Katsanis, 2009; Sheffield, 2010). The phenotype of BBS is indicative of defects in the function of cilia, and specifically in cilia-mediated signaling. Mutations in at least 12 genes (BBS1–BBS12) cause the syndrome (Jin and Nachury, 2009). Seven of the BBS gene products (BBS1, 2, 4, 5, 7, 8, and 9) form a biochemically stable complex, the BBSome (Nachury et al., 2007). The BBSome subunits are well conserved in organisms with cilia, indicating that the BBSome fulfills an important ciliary function. Recently, we identified Chlamydomonas reinhardtii mutants for BBS1, BBS4, and BBS7. Because large quantities of cilia can be isolated from C. reinhardtii, it was possible to carry out a direct biochemical comparison of wild-type and bbs mutant cilia. (Because cilia and flagella are essentially identical organelles, here, we refer to the two flagella of C. reinhardtii as cilia.) Remarkably, loss of the BBSome has little effect on the overall composition of cilia or the ciliary axoneme; rather, a small subset of membrane-associated proteins, several of which are predicted to have signaling function, are present at abnormal levels in bbs cilia (Lechtreck et al., 2009). A redistribution of ciliary membrane proteins is also characteristic for Bbs knockout mice (Berbari et al., 2008b; Domire et al., 2011; Seo et al., 2011; Zhang et al., 2011). However, the mechanism by which BBSome deficiency causes changes in ciliary protein composition remains unclear.

Data from Caenorhabditis elegans, C. reinhardtii, and mammalian cells have revealed that only intact BBSomes enter cilia (Blacque et al., 2006; Lechtreck et al., 2009; Seo et al., 2010, 2011). Within cilia, BBSomes are moved by intraflagellar transport (IFT). Previous work has shown that the membrane-associated signaling protein phospholipase D (PLD) accumulates abnormally in cilia of Chlamydomonas reinhardtii bbs mutants. Here we show that PLD is a component of wild-type cilia but is enriched ~150-fold in bbs4 cilia; this accumulation occurs progressively over time and results in altered ciliary lipid composition. When wild-type BBSomes were introduced into bbs cells, PLD was rapidly removed from the mutant cilia, indicating the presence of an efficient BBSome-dependent mechanism for exporting ciliary PLD. This export requires retrograde IFT. Importantly, entry of PLD into cilia is BBSome and IFT independent. Therefore, the BBSome is required only for the export phase of a process that continuously cycles PLD through cilia. Another protein, carbonic anhydrase 6, is initially imported normally into bbs4 cilia but lost with time, suggesting that its loss is a secondary effect of BBSome deficiency.

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Abbreviations used in this paper: AMPK, AMP-regulated protein kinase; BBS, Bardet-Biedl syndrome; CAH6, carbonic anhydrase 6; DIC, differential interference contrast; GPCR, G protein–coupled receptor; IFT, intraflagellar transport; MCHR1, melanin-concentrating hormone receptor 1; PA, phosphatidic acid; PE, phosphatidylethanolamine; PLD, phospholipase D; SSTR3, somatostatin receptor type 3.
transport (IFT), a process by which large membrane-associated protein particles are transported bidirectionally along the axonemal microtubules (Kozminski et al., 1993; Rosenbaum and Witman, 2002; Blacque et al., 2004; Lechtreck et al., 2009). These particles consist of over 20 different proteins organized into two sub-complexes, termed IFT complexes A and B (Cole and Snell, 2009). Kinesin motors move these particles toward the ciliary tip (anterograde IFT); cytoplasmic dynein 1b (cytoplasmic dynein 2 in mammals) returns the particles to the cell body (retrograde IFT). Ciliary assembly is largely unaffected in *C. reinhardtii* *bbs* mutants and in *Bbs* knockout mice, and organization of IFT complexes A and B appears unaffected in the *C. reinhardtii* mutants (Mykytyn et al., 2004; Lechtreck et al., 2009). BBS proteins are significantly less abundant than IFT proteins in cilia of *C. reinhardtii*, and only a subset of IFT particles carries BBSomes (Lechtreck et al., 2009). This suggests that the BBSome is an IFT cargo or cargo adapter dispensable for ciliary assembly and basic IFT in most cell types.

In mammalian cells, BBS deficiencies affect the presence of ciliary transmembrane proteins, specifically of G-protein–coupled receptors (GPCRs), incilia. Melanin-concentrating hormone receptor 1 (MCHR1) and somatostatin receptor type 3 (SSTR3) fail to localize to neuronal cilia in *Bbs2<sup><sub>-/-</sub></sup> and *Bbs4<sup><sub>-/-</sub></sup>* mice (Berbari et al., 2008a). The BBSome interacts with the ciliary targeting motif in the IP3 loop of SSTR3, and IP3<sub>SSTR3</sub>-GFP fusion proteins translocate into cilia in a BBSome–dependent manner (Berbari et al., 2008a; Jin et al., 2010; Domire et al., 2011). The BBSome could facilitate the transport of proteins from the plasma membrane through the barrier of the ciliary transition zone into the ciliary membrane proper (Nachury et al., 2010). However, recent data show that the localization of some ciliary GPCRs is unaffected by a BBSome deficiency, whereas still others, e.g., dopamine receptor 1 in *Bbs4<sup><sub>-/-</sub></sup>* mice and MCHR1 and the hedgehog effector Smoothened in *Bbs3<sup><sub>-/-</sub></sup>* mice, accumulate in BBSome-deficient cilia (Domire et al., 2011; Seo et al., 2011; Zhang et al., 2011). This suggests that both import and export of ciliary GPCRs are affected in *Bbs* mutants.

We previously identified several putative signaling proteins (phospholipase D [PLD], an AMP-regulated protein kinase [AMPK], and a single domain globin [THB1]) that accumulate excessively in the ciliary membranes of *C. reinhardtii* *bbs1, bbs4, and bbs7* mutants (Lechtreck et al., 2009). Although lacking receptor functions, these membrane-associated proteins could modulate ciliary signaling, e.g., by protein phosphorylation and the synthesis of signaling lipids, providing a potential explanation for the disruption of phototactic behavior that is a hallmark of *bbs* mutants in *C. reinhardtii*.

To investigate the mechanism by which these signaling proteins are accumulated in the *bbs* cilia, we performed a detailed analysis of one of the proteins, PLD, in wild type versus the *bbs4-1* mutant that is null for BBS4 (Lechtreck et al., 2009). We chose to focus on PLD, both because it has a mammalian orthologue, PLD6, and because of its likely involvement in phospholipid signaling (Munnik et al., 2000). We have found that BBSome deficiency causes a massive redistribution of PLD from the cell body to the ciliary membrane, that the biochemical defects in *bbs* cilia increase with time but can be rapidly corrected when wild-type BBSomes are introduced into the cytoplasm and cilia, that retrograde IFT acts upstream of the BBSome in the PLD export pathway, that PLD can enter the cilia independently of the BBSome and IFT, and, finally, that BBSome disruption causes secondary defects such as changes in the lipid composition of the ciliary membrane. We conclude that the *C. reinhardtii* BBSome functions downstream of IFT in the export phase of a process that cycles membrane signaling proteins through the cilia. The results further suggest that the absence of a protein from BBSome-deficient cilia is not necessarily due directly to failure of the BBSome to import the protein into the cilia.

**Results**

**PLD redistributes from the cell body into cilia in *bbs* mutants**

We previously reported, based on silver-stained gels, that PLD, AMPK, and THB1 are enriched in *C. reinhardtii* *bbs1, bbs4, and bbs7* cilia (Lechtreck et al., 2009). We have now generated an antibody that is specific for *C. reinhardtii* PLD (Fig. 1 A) and, using that antibody, are able to confirm that PLD is enriched in cilia of *bbs1, bbs4*, and *bbs7* mice (Fig. 1 B). In addition, we have extended the findings to a newly identified mutant for BBS8 (Fig. 1 B). This mutant has an insertion in the first exon of *BBS8* that results in a STOP codon predicted to terminate the normally 530-aa protein at aa 38 (Fig. S1 A). Therefore, this mutant is likely to be null for BBS8. Its isolated cilia lack BBS4, indicating that the BBSome without BBS8 is either unstable as in *bbs1-1* or is unable to enter the cilia as in *bbs7-1* (Lechtreck et al., 2009). To distinguish between these two possibilities, we assessed the level of BBS4 in the cytoplasm of the *bbs8-1* mutant; BBS4 was strongly reduced, indicating that the BBSome is largely unstable in the absence of BBS8 (Fig. S2 B). All four *bbs* mutants have normal length cilia (Fig. S2 A). Anterograde and retrograde IFT velocities as determined by differential interference contrast (DIC) microscopy are similar to that of wild type (Fig. S2 C); anterograde IFT frequency is also similar to wild type but retrograde IFT frequency appears to be slightly reduced (Fig. S2 D), although this was more difficult to measure because the retrograde particles are near the limit of detection for DIC. The levels of IFT particle proteins in the *bbs* cilia are similar to wild type (Fig. 1 C; and see Figs. 3 B and 6 A) or slightly elevated (Fig. S2 B and see Fig. 5 C); this slight increase, if real, may be an attempt by the mutant cells to compensate for loss of the BBSome. These results are consistent with our previous observation using immunofluorescence microscopy that the ciliary distribution of IFT complex A protein IFT139 and complex B protein IFT46 is unaffected in *bbs4-1* (Lechtreck et al., 2009).

Importantly, PLD can also be detected in wild-type cilia, particularly in the ciliary membrane, by means of Western blots probed with the anti-PLD antibody, indicating that PLD is a bona fide ciliary protein (Fig. 1, A and C). In wild type and *bbs4-1*, ciliary PLD is largely associated with the ciliary membrane, defined as the detergent phase of Triton X-114 phase partitioning (Fig. 1 C). To estimate just how great the...
In the mutant cilia, we used Western blotting to quantify the relative amounts of PLD in isolated membranes of wild-type and bbs4 mutant cilia. We determined that PLD is ~150–200x more abundant in the mutant versus wild-type ciliary membrane (Fig. 1 D). A similar enrichment was observed in cilia of mutant versus wild-type gametes (unpublished data).

To distinguish whether the observed enrichment of PLD in the cilia of bbs mutants is a result of a general increase in PLD levels in the mutant cells or to defects in ciliary trafficking of PLD, we determined the distribution of PLD in wild-type and bbs mutant cells and cilia. Western blotting revealed that the total amount of PLD in bbs4-1 cells actually was reduced by ~40% in comparison to wild-type cells (Fig. 1, E and H). Moreover, in wild-type, ~0.2% of the total PLD is in the cilia, whereas in the mutant, ~60% of the total PLD is in the cilia.

Immunofluorescence microscopy confirmed our biochemical data: PLD was readily detected in a spotted distribution along the length of bbs4-1 mutant cilia but was barely detectable in wild-type cilia (Fig. 1 F and see Fig. 4 B). Within mutant cilia the signal strength was more pronounced in the proximal region of the cilium. Labeling by the anti-PLD antibody was similar in wild-type and bbs4-1 cell bodies, albeit signal strength was reduced in the mutant. Label was distributed in a spotted pattern at the cell periphery (Fig. 1 G for bbs4-1); this could represent vesicles underlying the plasma membrane. In summary, the bbs4-1 mutation causes a dramatic redistribution of PLD from the cell body into cilia (Fig. 1 H).
The defects in protein content of bbs4 cilia increase with time

To distinguish whether the abnormal amounts of PLD in bbs cilia are deposited there during initial ciliary assembly or accumulate there with time after ciliary formation, we took advantage of the ability of C. reinhardtii to fully assemble cilia de novo within ~60 min after amputation of the cilia by pH shock (Fig. 3 and Fig. S3 E). We simultaneously deciliated large populations of wild-type and bbs4-1 cells and allowed them to synchronously regenerate their cilia; we then isolated the recently formed cilia at time points ranging from 80 to 380 min after deciliation and compared them by SDS-PAGE and Western blotting (Fig. 3 A). At the loadings used, PLD was not detected in wild-type whole cilia (Fig. 3 B, top), but small amounts of PLD were consistently detected in isolated wild-type ciliary membranes, where the PLD was maintained at a relatively constant low level during the time period analyzed (Fig. 3 B, bottom). In contrast, PLD was detectable in the newly assembled mutant whole cilia at least as early as 80 min after deciliation; it then accumulated steadily over the course of several hours as the cilia aged (Fig. 3, B [top] and C). The data indicate that PLD accumulation in mutant cilia is a dynamic process that continues in full-length steady-state cilia. The amounts and kinetics of accumulation of several ciliary transmembrane proteins, specifically PKD2, FMG1, and the mastigoneme protein, were similar in wild-type and mutant cilia, suggesting that the transport of these proteins is unaffected by BBS4 deficiency.

In the course of the cell cycle, cilia are normally formed after division, at least 6 h before our synchronously grown cells were used for biochemical analyses. We here term such 6-hour-old cilia “standard cilia.” Comparison of regenerated and standard cilia from wild type and bbs4-1 using silver-stained gels revealed a protein of ~24 kD that was present in wild-type cilia and in newly regenerated bbs4 cilia but was significantly reduced in standard bbs4 cilia (Fig. 3 D). The protein was tentatively identified as carbonic anhydrase 6 (CAH6) based on identification of a single peptide by mass spectrometry (Fig. S1 B); the identification was confirmed using an antibody to C. reinhardtii CAH6 (Fig. 3 B). Quantitative analysis of age-sorted cilia showed that at 80 min after deciliation the mutant cilia contained ~50% of the CAH6 present in comparable wild-type samples (Fig. 3, B and E). Over the next five hours CAH6 was progressively lost from mutant cilia, whereas little reduction of CAH6 was observed in wild-type cilia (Fig. 3, B, D, and E). Because a substantial amount of CAH6 is present in newly formed mutant cilia, it is unlikely that BBSome defects inhibit CAH6 ciliary import. The data rather suggest that bbs cilia increasingly lose the ability to retain CAH6. Thus, CAH6 loss from bbs cilia appears to be caused indirectly by the BBSome defect.

Excess PLD is rapidly removed from bbs cilia when BBSomes are reintroduced

The progressive redistribution of PLD from the cell body into cilia of bbs mutants could be caused by a failure of PLD ciliary export. Therefore, to test directly whether BBSomes...
PLD export from cilia requires retrograde IFT

We previously showed that the BBSome cycles in association with IFT through cilia (Lechtreck et al., 2009). To determine if this association is important for BBSome function and normal export of PLD from the cilium, we analyzed the mutant strain dhc1b-2 (previously referred to as dhc1bts). This strain contains a hypomorphic allele of the gene encoding the heavy chain of dynein 1b, the motor for retrograde IFT (Lechtreck et al., 2009; Witman, 2012). At the permissive temperature (21°C) used for all experiments involving dhc1b-2 here, ~60% of the mutant cells assembled two cilia that were slightly shorter than those of wild type (Fig. S3, A and B). The velocity and frequency of retrograde IFT and the amounts of the retrograde dynein subunits DHC1b and D1bLIC were reduced in the dhc1b-2 cilia (Fig. 5 A and Fig. S2, C and D). The amount of BBS4 was elevated in dhc1b-2 cilia, as expected for a protein cycled through the cilium by IFT. However, despite the presence of BBS4/BBSomes, PLD was highly enriched in the mutant cilia (Fig. 5 A), indicating that the BBSome-dependent export of PLD from cilia requires the retrograde motor for IFT.

As an independent test of the involvement of IFT in export of PLD, we analyzed the mutant ift74-1, which expresses a truncated version of the IFT complex B protein IFT74 (Fig. 5 B; unpublished data). The truncated IFT74 is assembled into complex B and transported into the cilium as part of the IFT particle; the ciliary levels of the other complex B proteins are near normal, but the amount of IFT complex A is greatly reduced. Compared with wild-type cilia, BBS4 was reduced by ~65% and
PLD was strongly accumulated in *ift74-1* cilia (Fig. 5 B). The data suggest that entry of BBSomes into cilia requires IFT74 and/or IFT complex A and that the residual IFT and BBSomes in *ift74-1* cilia are insufficient to remove PLD from cilia. The result is consistent with a recent observation in *C. elegans* that a mutation in complex A protein IFT144 abrogates entry of BBSomes into cilia (Wei et al., 2012).

IFT complex A has an important role in retrograde IFT (Pedersen and Rosenbaum, 2008; Pedersen and Christensen, 2012). To more directly investigate if complex A has a role in BBSome transport and PLD export, we analyzed strains *fla15-1* and *fla17-1*, which carry mutations in the complex A proteins IFT144 and IFT139, respectively (Piperno et al., 1998; Iomini et al., 2009). The *fla15-1* strain has a point mutation that converts a highly conserved cysteine in IFT144 to an arginine.

The *fla17-1* strain has an in-frame deletion of three exons of the *IFT139* gene and produces a smaller-than-normal version of IFT139 (Fig. 5 C; Iomini et al., 2009). These *fla* mutants fail to assemble cilia at elevated temperatures. At the permissive temperature used here IFT is sufficient for ciliary assembly but the velocity and frequency of retrograde IFT is reduced and the cilia develop small bulges containing complex B components, indicative of a defect in returning cargo to the cell body (Iomini et al., 2009). Compared with wild type, BBS4 was highly enriched in cilia of *fla17-1* (Fig. 5 C), suggesting that the BBSome also is not returned normally to the mutant cell body. Despite the abundance of BBSomes, PLD also was highly enriched in *fla17-1* cilia (Fig. 5 C), indicating that PLD export from cilia is dependent on IFT complex A. In contrast, the amounts of BBS4 and PLD were near normal in
cilia of fla15-1 (Fig. 5 C). The data suggest that IFT139 is involved in retrograde transport of BBSomes and removal of PLD from cilia.

PLD can enter cilia independently of IFT

Our data suggest that retrograde IFT particles transport PLD in a BBSome-dependent manner. Because PLD enters cilia through a BBSome-independent process, we investigated if IFT is required for entry of PLD into cilia. We took advantage of the fla10-1 mutant, which carries a temperature-sensitive mutation in one of the heavy chains of the anterograde IFT motor kinesin-2, so that IFT can be rapidly turned off by a temperature shift (Kozminski et al., 1995). At the permissive temperature (22°C), fla10-1 cilia contained slightly elevated amounts of PLD, which can be explained by the somewhat reduced amounts of IFT complex A and BBSomes (as indicated by reduced amounts of IFT139 and BBS4, respectively) in the mutant cilia (Fig. 6 A). Cilia isolated from fla10-1 that had been incubated for 2.5 h at the nonpermissive temperature (32°C) were largely depleted of IFT81 (complex B) and IFT139 (complex A). This loss of IFT proteins at the restrictive temperature was accompanied by an increase in ciliary PLD; in contrast, the amounts of PLD in wild-type and bbs4-1 cilia were essentially unaffected by the temperature shift. To determine the temporal sequence of IFT depletion and PLD accumulation in fla10-1 cilia, we analyzed cilia isolated at various time points (1–6 h) after shifting fla10-1 cells to 32°C (Fig. 6 B). During the incubation time the number of mutant cells carrying one or two cilia decreased from >90% to ~70% and the mean length of the cilia that they did have decreased from ~11 to ~6 µm; the number and length of cilia of wild type was unaffected at 32°C (Fig. S3, C and D). Starting at ~2 h at 32°C, by which time the level of IFT proteins was considerably reduced, PLD began to accumulate in fla10-1 cilia. Importantly, PLD continued to accumulate in fla10-1 cilia after 4 h at 32°C when IFT proteins were nearly undetectable; after ~6 h at 32°C it reached a level approaching that of bbs mutants (>30× above the level before temperature shift; Fig. 6 C). The data indicate that PLD entry into cilia does not require IFT. Note that despite the depletion of IFT proteins, BBS4 remained present in fla10-1 cilia. The accumulation of PLD in the presence of BBS4 provides additional evidence that BBSomes alone are insufficient to control PLD levels in cilia in the absence of IFT.

Discussion

BBSome deficiency has been shown to alter the membrane protein composition of cilia in C. reinhardtii and mammals (Berbari et al., 2008b; Lechtreck et al., 2009; Domire et al., 2011; Seo et al., 2011; Zhang et al., 2011). We showed previously that BBSomes travel through cilia in association with a subset of IFT particles (Lechtreck et al., 2009). These observations raise the possibility that BBSomes contribute to IFT-mediated transport of ciliary membrane proteins. Here we analyzed the role of the BBSome and IFT in the ciliary transport of PLD, a membrane-associated protein that accumulates in cilia of C. reinhardtii bbs mutants. Our data reveal that PLD continuously enters the cilia in a BBSome- and IFT-independent manner but that export of PLD from the cilia is dependent on the BBSome, probably functioning as a cargo adaptor for IFT.
there is a massive redistribution of PLD from the cell body into the cilia, such that about half of the PLD is now contained in the cilia. Taking into account the relative volumes of the cell body (>500 µm³) and the cilium (<0.5 µm³; and ignoring possible differences in the fraction of each that is accessible to PLD), PLD in bbs4-1 cells is enriched ~1,000X in the two cilia relative to the cell body. Because this concentration occurs in bbs mutants, PLD must be able to move into the cilia in the absence of BBSomes. A similar great enrichment was observed in cilia of the conditional mutant fla10-1 after IFT proteins (but not BBS4) were depleted by incubation at the nonpermissive temperature. This strongly suggests that entry of PLD into cilia can occur even in the absence of IFT. Recently, Kee et al. (2012) reported that small molecules and proteins ≤40 kD can freely pass from the cytoplasm into cilia. Thus, PLD, which is only 25 kD, may enter the cilia slowly by diffusion and then become trapped there in the absence of an active export mechanism.

Because PLD accumulates progressively over time in BBSome- and IFT-deficient cilia, it is likely that the protein is continuously cycled through the cilium in wild-type cells, with IFT-dependent export mediated by the BBSome. The high efficiency of this PLD export pathway, as demonstrated by the rapid clearing of accumulated PLD when BBSomes are reintroduced into bbs mutant cilia, suggests that PLD entering wild-type cilia will be rapidly picked up and removed from the organelle. A system in which the movement of a protein into the cilium is BBSome independent but its export is BBSome dependent would allow fine tuning of the amount of the protein in the cilium by regulating the number of BBSomes that enter the cilium or by regulating the affinity of the BBSome for the protein.

A similar role for the BBSome in mammalian primary cilia is suggested by recent observations on dopamine receptor 1, a GPCR that is rapidly transported to and from cilia in response to environmental cues (Domire et al., 2011). Import of dopamine receptor 1 into cilia is BBSome independent whereas translocation of the protein from cilia requires the BBSome (Domire et al., 2011; Zhang et al., 2011). In addition, very recent results indicate that accumulation of the hedgehog signaling proteins Smoothened and Patched 1 occur in cilia of cells from Bbs⁻/⁻ mice (Zhang et al., 2012). Like PLD in C. reinhardtii, Smoothened is believed to continuously shuttle through the cillum (Ocbina and Anderson, 2008). Thus, BBSome-mediated export may be a feature of many membrane-associated proteins that undergo such cycling.

**BBSome function depends on retrograde IFT**

To investigate the role of IFT in the BBSome-dependent export of PLD from the cilium, we analyzed the distribution of BBS4 and PLD in cilia isolated from the dynein 1b mutant dhc1b-2, which is characterized by reduced retrograde movement of IFT particles. Using silver-stained 1D and 2D gels, we previously determined that THB1 is enriched in dhc1b-2 cilia, but we could not ascertain the levels of PLD in these cilia (Lechtreck et al., 2009). Now, using our anti-PLD antibody and Western blotting, we found that PLD also is significantly accumulated in dhc1b-2 cilia despite the presence of BBSomes. Likewise, BBSomes are present albeit reduced during PLD accumulation in fla10-1 cilia at restrictive temperature. Therefore, the presence of BBSomes in cilia is insufficient to remove PLD when IFT is defective. Because the primary effect of the dhc1b-2 mutation is on retrograde IFT, these results indicate that retrograde IFT is critical...
for PLD export from the cilium. Inasmuch as loss of the BBSome has little if any effect on IFT (Figs. 1 C, 3 B, 6 A, and S2; Lechtreck et al., 2009), but defects in retrograde IFT affect export of both the BBSome and PLD, IFT must function upstream of the BBSome in the PLD export pathway; i.e., retrograde IFT moves the BBSome out of the cilium, and the BBSome in turn carries PLD from the cilium (Fig. 5 D).

Export of Smoothened from mammalian primary cilia similarly is dependent on the retrograde IFT motor, as indicated by the accumulation of Smoothened in cilia of mouse cells having a mutation in the dynein 2 heavy chain Dync2h1 (Ochina and Anderson, 2008) or in which the same protein was knocked down by shRNA (Kim et al., 2009) or inhibited pharmacologically (Firestone et al., 2012). Thus, IFT-dependent BBSome-mediated export of signaling molecules from the cilium may be widespread. However, another hedgehog signaling protein, Gli2, which also is believed to cycle though the cilium (Kim et al., 2009), does not accumulate in Bbs6−/− mice (Zhang et al., 2012), suggesting the existence of distinct export (and import) mechanisms for different ciliary proteins.

**IFT complex A is critical for BBSome export**

The aforementioned observations suggest that coupling of PLD to retrograde IFT particles requires the BBSome. To explore how the BBSome interacts with the IFT machinery, we used mutants defective in IFT particle proteins. IFT complex A proteins and BBS4 are similarly decreased in cilia of ift74-1, which expresses a truncated complex B IFT74 protein. PLD is highly enriched in ift74-1 cilia, indicating that the residual BBSomes are insufficient for its export. This suggests that IFT74 is critical for the entry of IFT complex A into cilium and that complex A could mediate binding of the BBSome to IFT complex B and the associated dynein motors during retrograde IFT. To test for a possible role of IFT complex A as a docking site for the BBSome, we analyzed fla15 and fla17-1, which are defective in the IFT complex A proteins IFT144 and IFT139, respectively, and have reduced retrograde IFT (Piperno et al., 1998; Iomini et al., 2009). The distributions of BBS4 and PLD were nearly normal in fla15, whereas PLD and in particular BBS4 were enriched in cilia of fla17-1. Therefore, IFT139 could be involved in binding the BBSome to IFT complex A during retrograde IFT, rendering it critical for the export of the BBSome and PLD from cilia (Fig. 5 D). It may be relevant that the Tubby family protein TULP3 similarly interacts with complex A in mammals to promote ciliary trafficking of a subset of GPCRs, including MCHR1, and that retrograde transport of TULP3 specifically requires IFT139 (Mukhopadhyay et al., 2010).

In summary, PLD accumulates in cilia of mutants with defective BBSomes (bbs4-1, bbs1-1, bbs7-1, and bbs8-1; Lechtreck et al., 2009), with reduced amounts of BBSomes in cilia (ift74-1), and with reduced retrograde IFT of the BBSome (dhc1b-2 and fla17-1). We propose that the BBSome functions as a cargo adapter linking PLD to retrograde IFT particles to ensure export of PLD from cilia. Further work will be necessary to understand how PLD is delivered to the cilium, how PLD interacts with the BBSome during the export phase of its cycle, and how the BBSome interacts with IFT complexes A and B.

**BBSome function in IFT assembly and turnaround**

In *C. elegans* mutants null for bbs-7 and bbs-8, IFT complex A separates from complex B and the two complexes are transported anterogradely by kinesin-2 and OSM-3, respectively (Ou et al., 2005). These observations led to the hypothesis that the BBSome couples complex A to complex B and coordinates the activity of the two anterograde IFT motors. However, more recently it has been reported that a hypomorphic mutation at a
conserved site in the complex A protein DYF-2 (the *C. elegans* homologue of IFT144) causes complete dissociation between the BBSome and moving IFT particles, yet complexes A and B remain associated in anterograde IFT (Wei et al., 2012). These results indicate that in *C. elegans* the BBSome is not essential for stabilizing the binding of complex A to complex B, in good agreement with our previous observation that the colocalization of complexes A and B is not affected in cilia of *C. reinhardtii bbs4-1* mutants (Lechtreck et al., 2009). In the *C. elegans* IFT144 mutant, the BBSome is still formed but remains at the base of the cilium. Moreover, complex B but not complex A accumulates at the tip of the cilium, suggesting that complex B fails to be coupled to complex A for retrograde IFT. Based on these observations, Wei et al. (2012) proposed that in wild-type *C. elegans* the BBSome regulates assembly of complexes A and B into intact IFT particles at the base of the cilium, and is then carried as cargo by IFT to the tip of the cilium, where it has a similar role in reorganizing the IFT particle for retrograde transport. In the mutant, the BBSome still functions to assemble the IFT particle at the base of the cilium, but because it is no longer transported to the ciliary tip, IFT particle reorganization is defective and complex B accumulates there.

We previously reported that in *C. reinhardtii* the BBSome is not an integral part of the IFT machinery but rather appears to be a cargo or cargo adapter, a conclusion now supported by the observations of Wei et al. (2012). However, in *C. reinhardtii* loss of the BBSome does not seem to affect the assembly of complexes A and B (Lechtreck et al., 2009), does not lead to an accumulation of complex B at the tip of the cilium (Lechtreck et al., 2009), and does not obviously alter the balance of complex A to complex B within the cilium as would be expected if complex B were being left behind during retrograde transport (Figs. 5 C, 6 A, and S2 B). Therefore, in contrast to the situation in *C. elegans*, it is unlikely that the BBSome has a key role in regulating IFT particle assembly at either the base or the tip of the *C. reinhardtii* cilium.

The function of PLD in *C. reinhardtii*

The function of PLD in *C. reinhardtii* is likely to involve phospholipid signaling (Meijer et al., 2002). Consistent with this, we found that the accumulation of PLD in cilia of *C. reinhardtii* mutants is correlated with major changes in the phospholipid content of the ciliary membrane. It is possible that PLD cycles through the wild-type cilium as part of a system that monitors and responds to the cell’s environment (e.g., to light or to osmotic conditions) via a phospholipid-based signaling pathway. We previously reported that *C. reinhardtii bbs* mutants are nonphototactic. Phototaxis is dependent on membrane depolarization and Ca\(^{2+}\) influx (Witman, 1993), both of which activate PLD in *C. reinhardtii* (Munnik et al., 2000). Changes in membrane lipid composition could affect the functioning of membrane channels and hence phototaxis. Consistent with this, we previously observed that *bbs4-1* cells with newly formed cilia were able to phototax for over an hour, but lost that ability as the cilia aged and presumably accumulated PLD and acquired an altered membrane lipid composition (Lechtreck et al., 2009).

*C. reinhardtii* PLD has homologues in mammals and many other ciliated eukaryotes

Our finding that *C. reinhardtii* PLD is apparently a bona fide ciliary protein that cycles through the cilium raises the question of whether a homologue occurs in the cilia of other organisms. Close homologues of *C. reinhardtii* PLD are encoded in the genomes of many vertebrates, nonvertebrate metazoans (with *C. elegans* being a notable exception), and some ciliated protists; they are absent from *Arabidopsis thaliana*, yeasts, and other nonciliated organisms (Fig. S4, A and B). Such a phylogenetic distribution is often indicative of a centriolar or ciliary function of the gene product. The mammalian homologue of PLD is PLD6 (*C. reinhardtii* to *H. sapiens* BLAST E = 8e−37; 40% identity); in a phylogenetic analysis PLD6 groups more closely with *C. reinhardtii* PLD than with other murine PLDs (Fig. S4 B). In mice, overexpressed PLD6 localizes to mitochondria and is anchored to the mitochondrial surface by a transmembrane domain (Huang et al., 2011). This transmembrane domain is not predicted in *C. reinhardtii* PLD but, like *C. reinhardtii* AMPK and CAH6, the protein is predicted to be myristoylated and palmitoylated (Fig. S4 C), likely accounting for its association with the ciliary membrane. PLD6 is widely expressed in human tissues; in mice, expression is especially high in lung and testis, tissues that are rich in cilia, and *Plde−/−* mice are infertile (Huang et al., 2011; Watanabe et al., 2011). It will be of interest to determine if mammalian PLD6 functions also in cilia, and whether BBSome deficiency promotes ciliary accumulation of PLD6. It may be relevant that cilia of *Bbs−/−* mice accumulate unusual intraciliary vesicles at the ciliary tip (Shah et al., 2008); this might be brought about by an excess of ciliary DAG caused by ciliary accumulation of PLD6. Phospholipid signaling also has been implicated in the functioning of olfactory cilia (Klasen et al., 2010), and malfunction of olfactory cilia has been reported in *Bbs8−/−* mice (Tadenev et al., 2011).

BBSome deficiency results in secondary defects in ciliary composition

Quantitative analysis of age-sorted cilia revealed that the ciliary protein CAH6 is progressively lost from *bbs4-1* cilia. Because CAH6 is initially present in mutant cilia, it is unlikely that BBSome defects inhibit CAH6 ciliary import. Rather, the progressive loss of CAH6 from *bbs4-1* cilia is likely to be caused indirectly by BBSome deficiency. Loss of a peripheral membrane protein such as CAH6 could follow from the severe changes in membrane lipid content that occur in *bbs4-1* cilia, presumably as a result of increased ciliary PLD activity; the ability of proteins, such as CAH6, which is predicted to be fatty acid modified (Fig. S4 C), to associate with membranes is influenced by the lipid composition of the membrane (Levental et al., 2010). An altered lipid composition of *bbs* cilia also could contribute in other ways to changes in the pattern of ciliary membrane proteins. Loss of certain GPCRs from cilia has been reported in murine *Bbs−/−* mutants (Berbari et al., 2008b). In light of our data on CAH6 in *bbs* mutant cilia of *C. reinhardtii*, it seems possible that absence of GPCRs from mammalian cilia also could be an indirect consequence of the loss of the BBSome.
BBS likely involves progressive deterioration of ciliary function

The observation that ciliary defects resulting from BBSsome deficiency increase with time potentially has implications for the pathomechanism of BBS in mammals. Several features of BBS in murine Bbs−/− models or human patients with BBS also increase with time or manifest themselves only years or decades after birth (Beales, 2005). The outer segments of photoreceptor cells in the eye, for example, initially develop and function normally in Bbs2 and Bbs4 knockout mice. About 6 wk postnatally, rhodopsin mislocalization becomes apparent and retinal degeneration begins (Nishimura et al., 2004). Therefore, BBSsome defects appear to initiate a cascade of events that increasingly deteriorate ciliary protein composition and function. The time scales in which ciliary defects develop in different organisms and different tissues within an organism may be different. Nevertheless, the mechanism by which cells use the BBSsome to control ciliary protein transport and to maintain the balance of ciliary signaling proteins might very well be conserved.

Materials and methods

Strains and culture conditions

Wild-type and mutant strains of C. reinhardtii were mostly maintained as described in Lechtreck et al. (2009). The bbs8-1 strain was generated as described in the legend of Fig. S1. The retrograde IFT mutants fla15 and fla17-1 were aerated without CO2 supplement and maintained at 22°C.

Antibodies and Western blotting

Using primers 5′-CGCGAATTCATGGGTTGCGCCAGCTCC-3′ and 5′-CGC-GAATTCATGGGTTGCGCCAGCTCC-3′ and 5′-CGC-GAATTCATGGGTTGCGCCAGCTCC-3′, a partial cDNA encoding C. reinhardtii PLD was obtained by PCR and cloned into bacterial expression vectors pGEX and pMal. Truncated PLD fused to maltose-binding protein was used as an antigen to immunize two rabbits, and a fusion protein of truncated PLD and glutathione S-transferase was used to affinity purify the antiserum produced by GenScript. Anti-PLD was diluted 1:2,000 for Western blotting and 1:100 for immunofluorescence microscopy. Polyclonal antibodies to C. reinhardtii CAH6 (1:2,000) and FAP12 (1:1,000) were provided by J.V. Moroney (University of Muenster, Muenster, Germany), and for temperature-shift experiments with fla10-1, cells were concentrated and incubated for 10 min at 35°C, 37°C, or 38°C in an incubator. For time course experiments, equal aliquots of concentrated samples were incubated in the dark for 10 min at room temperature (22°C) or at 32°C in an incubator. For deciliation by pH shock, 0.5 M acetic acid was added to cells in M medium with rapid stirring until the pH reached 4.2; after 50 s the solution was neutralized using 0.25 N KOH. Cells were immediately placed into fresh culture medium and allowed to regenerate cilia in bright light at 37°C with shaking (60 rpm). Isolated flagella were examined using transmission electron microscopy (EM) to verify that the axonemes were free of membranes and matrix fragments.

Isolation of cilia

Flagellar isolation using the dibucaine method and flagellar amputation by pH shock were performed as previously described by Witman (1986) and Lefebvre (1995), respectively. In brief, cells were concentrated and repeatedly washed in 10 mM HEPES, pH 7.4, resuspended in HMS (10 mM HEPES, pH 7.4, 5 mM MgSO4, and 4% wt/vol sucrose), and placed on ice; all subsequent steps were performed at 4°C. 2 ml of dibucaine (25 mM in H2O) were added per 10 ml of sample; cells were rapidly drawing them up and down in a pipette. 20 ml of 0.7 mM EGTA in HMS was then added to the suspension. Cell bodies were removed by differential centrifugation (3 min at 1,150 g), the supernatant was collected, and the remaining cell bodies were removed by centrifugation (10 min at 1,700 g) through a sucrose cushion (10 ml 25% sucrose in HSM). The supernatant was harvested and cilia were pelleted at 27,000 g for 20 min. For deciliation by pH shock, 0.5 M acetic acid was added to cells in M medium with rapid stirring until the pH reached 4.2; after 50 s the solution was neutralized using 0.25 N KOH. Cells were immediately placed into fresh culture medium and allowed to regenerate cilia in bright light at 37°C with shaking (60 rpm). Isolated flagella were examined using transmission electron microscopy (EM) to verify that the axonemes were free of membranes and matrix fragments.

Lipidomic and metabolomic analyses of cilia

We used two distinct approaches for the lipidomic analysis of cilia, liquid chromatography to mass spectrometry and an approach based solely on mass spectrometry. In brief, cilia were extracted with methanol/chloroform and lipid extracts were separated by gradient elution normal phase HPLC and the analytes detected by positive atmospheric pressure chemical ionization. For metabolomic analysis by gas chromatography phase HPLC and the analytes detected by positive atmospheric pressure chemical ionization. For metabolomic analysis by gas chromatography/mass spectrometry, samples were derivatized to their trimethylsilyl derivatives. Chromatography was on a 30-M DB-5MS column. Full scan
electron ionization mass spectrometry was performed using a Quattro II mass spectrometer (Waters). The accuracy of quantitation is ±10%.

For quantitative analysis of phospholipids, ~100 µg of cilia (wet weight) from each sample was subjected to lipid extraction. In brief, the sample was dissolved in 200 µl of liquid chromatography grade water. Samples were spiked with 10 µl of a mixture of internal standards containing 20 pmol DAG 17:0–17:2, 24 pmol PA 17:0–17:0, 52 pmol PE 17:0–17:0, 7.5 pmol phosphatidylglycerol (PG) 17:0–17:0, 43 pmol phosphatidylserine (PS) 17:0–17:0, 40 pmol phosphatidylcholine (PC) 18:3–18:3, 54 pmol phosphatidylinositol (PI) 17:0–17:0, 20 pmol ceramide [Cer] 18:1/2/17:0, 0.40 pmol sphingomyelin (SM) 18:1/2/17:0, 20 pmol galactosylceramide [GalCer] 18:1/2/12:0, and 20 pmol lactosylceramide [LaCer] 18:1/2/12:0. 265 µl of methanol was added to the mixture and agitated for 10 min for homogenization. Then, 730 µl of chloroform were added and the mixture was agitated for an additional 1 h. The lower organic phase was collected and evaporated in a Speedvac concentrator at room temperature. Lipid extracts were dissolved in 100 µl of 0.1% methylamine in methanol and subjected to quantitative lipid analysis on a hybrid QSTAR Pulsa quadrupole time-of-flight mass spectrometer (MDS Sciex). Samples were infused with a TriVersa NanoMate robotic nanoflow ion source (Advion Biosciences, Inc.). DAG, PA, PS, PE, PI, and PG species were quantified by negative ion mode multiple precursor ion scanning analysis (Ejsing et al., 2006).

Miscellaneous
For mass spectrometric identification of CAH6, bands of interest were excised from silver-stained gels and digested overnight with trypsin, and the eluted peptides were analyzed using a matrix-assisted laser desorption/ionization time-of-flight mass spectrometer (Kratos Axima QIT; Shimadzu Instruments). Peptides were analyzed in positive ion mode using a nanoflow ion source (Advion Biosciences, Inc.). DAG, PA, PS, PE, PI, and PG species were quantified by negative ion mode multiple precursor ion scanning analysis (Ejsing et al., 2006).

Online supplemental material
Fig. S1 A shows a molecular map of bbs8-1; the sequence of CAH6 is depicted in Fig. S1 B. Fig. S2 analyzes ciliary length and IFT in dhtc1b-1 and the bbs mutant strains and shows protein content of wild-type and bbs mutant cilia and cell bodies. Fig. S3 shows ciliary numbers and length for fla10-1 and dhtc1b-1 and compares ciliary regeneration in wild type and mutant strains. For mass spectrometric analysis, cilia were isolated from cell bodies using a protease cocktail, lysed, and analyzed by mass spectrometry. For mass spectrometric analysis, cilia were isolated from cell bodies using a protease cocktail, lysed, and analyzed by mass spectrometry (MDS Sciex). Samples were infused with a TriVersa NanoMate robotic nanoflow ion source (Advion Biosciences, Inc.). Peptides were analyzed in positive ion mode using a nanoflow ion source (Advion Biosciences, Inc.). DAG, PA, PS, PE, PI, and PG species were quantified by negative ion mode multiple precursor ion scanning analysis (Ejsing et al., 2006).

Identification of ciliary localization sequences within the third intracellular loop of G protein-coupled receptors was performed using a matrix-assisted laser desorption/ionization time-of-flight mass spectrometer (Kratos Axima QIT; Shimadzu Instruments). Peptides were analyzed in positive ion mode using a nanoflow ion source (Advion Biosciences, Inc.). DAG, PA, PS, PE, PI, and PG species were quantified by negative ion mode multiple precursor ion scanning analysis (Ejsing et al., 2006).

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References


Figure S1. **Molecular characterization of an inser- tional mutation in *C. reinhardtii* BBS8 and sequence of CAH6.** (A) Molecular characterization of an inser- tional mutation in *C. reinhardtii* BBS8. Wild-type *C. reinhardtii* was transformed with a promoter-less fragment of the AphVII gene conferring resistance to hygromycin (Berthold et al., 2002). Transformants were screened for lack of phototaxis; the nonphototactic strain LP20.2 was selected and analyzed by re- striction enzyme site-directed amplification PCR (González-Ballester et al., 2005; Brown et al., 2012). The strain has an insertion in exon 1 codon 27 be- tween basepairs 82 and 83 (indicated by the triangle; counting from the ATG); no deletion of genomic DNA was detected. This insertion causes an in-frame STOP after 38 residues/codons. The arrows represent the flanking DNA amplified by PCR. The third base of the inserted promoter-less AphVII was altered from T to A. We designated this allele bbs8-1 and here refer to the strain carrying it by the same name. (B) Sequence of CAH6. The peptide identified by mass spectrometry is shown in red.
Figure S2. Analysis of ciliary length and IFT in dhc1b-2 and bbs mutants. (A) Ciliary length of wild type and the four bbs mutant strains. (B) Western blot analysis of cilia isolated from wild type and bbs4-1. For quantification of relative protein amounts, values for band intensities were adjusted for IC2 loading and the ratios (left, parentheses) were then calculated. Velocity (C) and frequency (D) of anterograde and retrograde IFT for wild type and dhc1b-2 and bbs mutants. Standard deviations are indicated in A, C, and D; t-test probability data are shown in D.
Figure S3. Cilia length and number as a function of time and temperature in dhc1b-2, fla10-1, and bbs4-1 mutants. (A and B) Ciliary number and ciliary length, respectively, for the dhc1b-2 mutant at the permissive temperature (21°C). (C) Ciliary number for wild-type and fla10-1 cells before (0 h) and at various time points (1–6 h) after shifting the cells to 32°C. The data shown are from a single representative experiment out of two repeats. The number of cilia of wild-type or fla10-1 cells maintained at 22°C remained at the 0-h level throughout the time course (not depicted). (D) Ciliary length of wild-type and fla10-1 cells before (0 h) and after various times (1–6 h) of incubation at 22 or 32°C. (E) Ciliary length at various time points after deciliation by pH shock. Measurements are based on isolated cilia viewed by dark-field microscopy. Red, bbs4-1; gray, wild type; dashed lines, length of wild-type cilia (black, 10.8 ± 0.91 µm, n = 29) and bbs4-1 cilia (red, 11.8 ± 1.18 µm, n = 26) before deciliation. At least 24 cilia from different cells were measured for each time point. Standard deviations are indicated in B, D, and E.
Figure S4. **Alignment and phylogenetic analysis of PLDs.** (A) Alignment of *C. reinhardtii* and vertebrate PLDs. #, active site; ^, signature sites, #, both.

(B) Rooted phylogenetic tree of murine PLDs (Mm_PLD1 [NCBI Protein accession no. NP_001157528.1], Mm_PLD2 [GenBank accession no. AAH38563]), Mm_PLD3 [GenBank accession no. AAH38563]), Mm_PLD4 [GenBank accession no. AAH38563]), Mm_PLD5 [GenBank accession no. AAH62849], and Mm_PLD6 [GenBank accession no. CAI24298.1]), Arabidopsis thaliana PLDs (AtPLDalpha1 [GenBank accession no. AEE75720]), AtPLDdelta [GenBank accession no. AED93432]), Drosophila melanogaster zucchini (GenBank accession no. AAM49862.1), Tetrahymena thermophila TTHERM_02188720 (NCBI Protein accession no. XP_001028688), and *C. reinhardtii* PLD (NCBI Protein accession no. XP_001693080). The tree is based on a CLUSTALW multiple sequence alignment using default settings.

(C) N-Terminal sequences of PLD, CAH6, and AMPK. Residues predicted to be myristoylated are shown in red and residues predicted to be palmitoylated in green. Online tools (Myristylator, Myr Predictor, and CSS-Palm 2.0) were used for prediction of fatty acid modification.