Single-particle tracking localization microscopy reveals nonaxonemal dynamics of intraflagellar transport proteins at the base of mammalian primary cilia

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ABSTRACT Primary cilia play a vital role in cellular sensing and signaling. An essential component of ciliogenesis is intraflagellar transport (IFT), which is involved in IFT protein recruitment, axonemal engagement of IFT protein complexes, and so on. The mechanistic understanding of these processes at the ciliary base was largely missing, because it is challenging to observe the motion of IFT proteins in this crowded region using conventional microscopy. Here, we report short-trajectory tracking of IFT proteins at the base of mammalian primary cilia by optimizing single-particle tracking photoactivated localization microscopy for IFT88-mEOS4b in live human retinal epithelial cells. Intriguingly, we found that mobile IFT proteins "switched gears" multiple times from the distal appendages (DAPs) to the ciliary compartment (CC), moving slowly in the DAPs, relatively fast in the proximal transition zone (TZ), slowly again in the distal TZ, and then much faster in the CC. They could travel through the space between the DAPs and the axoneme without following DAP structures. We further revealed that BBS2 and IFT88 were highly populated at the distal TZ, a potential assembly site. Together, our live-cell single-particle tracking revealed region-dependent slowdown of IFT proteins at the ciliary base, shedding light on staged control of ciliary homeostasis.

INTRODUCTION

The primary cilium is an essential organelle responsible for multiple sensory and signaling activities (Goetz and Anderson, 2010). It originates from the mother centriole–derived basal body, with distal appendages (DAPs) at the periphery of the centriolar distal end and the transition zone (TZ), slowly again in the distal TZ, and then much faster in the CC. They could travel through the space between the DAPs and the axoneme without following DAP structures. We further revealed that BBS2 and IFT88 were highly populated at the distal TZ, a potential assembly site. Together, our live-cell single-particle tracking revealed region-dependent slowdown of IFT proteins at the ciliary base, shedding light on staged control of ciliary homeostasis.

To perform their functions, IFT proteins must migrate from the DAPs, the recruiting site of IFT proteins at the base of mammalian primary cilia by optimizing single-particle tracking photoactivated localization microscopy for IFT88-mEOS4b in live human retinal epithelial cells. Intriguingly, we found that mobile IFT proteins "switched gears" multiple times from the distal appendages (DAPs) to the ciliary compartment (CC), moving slowly in the DAPs, relatively fast in the proximal transition zone (TZ), slowly again in the distal TZ, and then much faster in the CC. They could travel through the space between the DAPs and the axoneme without following DAP structures. We further revealed that BBS2 and IFT88 were highly populated at the distal TZ, a potential assembly site. Together, our live-cell single-particle tracking revealed region-dependent slowdown of IFT proteins at the ciliary base, shedding light on staged control of ciliary homeostasis.

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is essential for molecular transport along the ciliary axoneme (Taschner et al., 2012). Once ciliogenesis is initiated, IFT proteins assemble with molecular motors and adapter proteins such as BBSome molecules, proteins of the Bardet–Biedl syndrome (BBS)-associated complex, which is important for cilium biogenesis, to facilitate precursor transport (Rosenbaum and Witman, 2002; Ou et al., 2005). Although IFT proteins such as IFT52 are recruited to the DAPs, it is unclear whether this location is the site where the different IFT proteins assemble and where BBS proteins, molecular motors, and cargoes integrate with IFT proteins (Rosenbaum and Witman, 2002; Pedersen et al., 2006; Wren et al., 2013; Craft et al., 2015). All of these proteins may be recruited to the DAPs and assembled there, or recruited there but assembled somewhere else, or individually recruited at different locations. One major missing link in the IFT process is how IFT proteins move between the DAPs and the ciliary axoneme. At the macromolecular scale, the ~130-nm distance between the tips of DAPs and the axoneme is relatively large. It is unclear whether DAP-recruited IFT proteins diffuse across the gap to the axoneme or whether they travel along the DAP structures.

Kymograph analysis has enabled particle tracking of anterograde and retrograde IFT dynamics along ciliary and flagellar axonemes (Snow et al., 2004; Engel et al., 2009; Wren et al., 2013; Craft et al., 2015). When it was integrated with total internal reflection (TIRF) microscopy and fluorescence recovery after photobleaching, signal-to-noise ratio (SNR) was much improved, and thus the mechanisms of IFT regulation along the cilia and flagella were further understood (Engel et al., 2009; Wren et al., 2013; Craft et al., 2015). Kymograph studies have shown that IFT proteins move more slowly toward the base (Prevo et al., 2015), but when examining the kymographs closely, one frequently sees an ultrabright “band” of IFT proteins present at the ciliary base (e.g., see Ye et al., 2013), reflecting the ultrahigh density of IFT proteins packed in the tiny volume surrounding the DAPs and TZ at the ciliary base. Advanced imaging techniques have been implemented to track single molecules in flagella and cilia, including Fourier filtering to achieve single-molecule sensitivity for motors and IFT proteins (Prevo et al., 2015; Mjalkovic et al., 2017), low-concentration expression to track surface proteins SSTR3 and Smo (Ye et al., 2013), and SNAP-tag labeling to track sporadic single-molecule events of Smo (Milenkovic et al., 2015). However, none of these techniques addressed IFT protein dynamics at the highly dense area of the ciliary base.

Superresolution microscopy such as direct stochastic optical reconstruction microscopy (dSTORM) can be used to localize molecules at a high precision for fixed-cell imaging (Heilemann et al., 2008). Single-particle tracking photoactivated localization microscopy (sptPALM) is a live-cell localization imaging technique for tracking a photoactivated subpopulation of fluorescent proteins, each emitting photons for a finite number of frames (Hess et al., 2007; Manley et al., 2008; Frost et al., 2010; Hoze et al., 2012; Rossier et al., 2012; Izeddin et al., 2014). It has been applied to study the dynamics of high-density molecules, including AMPA receptors, integrins, transcription factors, and actin assembly (Frost et al., 2010; Hoze et al., 2012; Rossier et al., 2012; Izeddin et al., 2014). Recently, sptPALM has been applied to reveal the dynamic interactions of IFT proteins in the ciliary compartment of Caenorhabditis elegans cilia (Oswald et al., 2018). Diffusion analysis and distribution analysis of sptPALM data have been performed to reveal novel dynamic characteristics of biomolecules (Manley et al., 2008; Meijering et al., 2012; Persson et al., 2013; Izeddin et al., 2014). The challenge of using sptPALM to address specific biological problems is that every problem has unique characteristics such as molecular density, background noise, and speed of motion that require optimization of the sptPALM protocol.

Many important findings concerning IFT dynamics have been revealed by TIRF microscopy, which enables tracking of bright particles in flagella or cilia (e.g., see Engel et al., 2009; Wren et al., 2013; Craft et al., 2015). sptPALM, on the other hand, is a single molecule–based assay, allowing short-distance tracking of single fluorophores when the particles are crowded or the fluorophores are normally hidden among other signals. For example, one may be able to observe motion of individual IFT proteins shown as the background fluorescence along the ciliary compartment, which is dimmer than that of moving IFT trains, or one may be able to see IFT dynamics in highly crowded regions such as the ciliary base and tip. Thus, sptPALM serves as a good complementary technique for TIRF microscopy for dynamic studies of IFT proteins.

In this study, we revealed novel IFT protein dynamics at the ciliary base by optimizing the sptPALM technique. With this unprecedented imaging capability for ciliary studies, we uncovered possible structural hindrance and preferred orientations along the paths IFT proteins travel, elucidating the mechanisms of different ciliogenesis-associated processes of IFT proteins, including protein assembly, microtubule track engagement, and passage gating.

RESULTS

sptPALM enabled dynamic studies of IFT proteins

We developed an sptPALM-based imaging protocol using photoconvertible fluorescent proteins to explore the dynamics of IFT proteins at the base of primary cilia. The excessive crowding of IFT proteins in this region and their high velocities necessitated multiparameter optimization of sptPALM to balance SNR, frame rate, trajectory duration, phototoxicity, and emission isolation. Human retinal pigment epithelial cells (hTERT-RPE-1; hereafter RPE-1) with stably expressed IFT88 fused to mEos4b, a monomeric photoconvertible protein (Paez-Segala et al., 2015), were created. The cell morphology, average ciliary length, and ciliation frequency of this cell line were similar to those of wild-type RPE-1 cells. Single-molecule detection was achieved by implementing 405-nm photoconversion with its power being gradually and carefully incremented to photoconvert a finite population of IFT88-mEos4b fluorophores that could be optically isolated in this highly crowded region. These photoconverted fluorophores were then excited by a 561-nm laser (Figure 1A and Supplemental Movie S1). The laser power was optimized such that it had to be strong enough to reach an acceptable SNR while weak enough to extend the imaging duration to be as long as possible and weak enough to minimize phototoxicity. Similar to IFT88 in wild-type RPE-1 cells, IFT88-mEos4b also formed a high-intensity punctum at the ciliary base (Figure 1B). In contrast to most other sptPALM-based studies, which collect data from a cellular or subcellular volume of ~1000 μm3, our imaging volume was usually confined to less than 1 μm3, and thus the number of emitting events in each experimental run was very limited. Each of the photoconverted mEos4b emitting events was recorded as a short trajectory of IFT88 movement lasting several to tens of time points (Figure 1A). Imaging at 30 ms per frame was adopted to collect enough photons for single-molecule fitting while being fast enough to trace rapid diffusing movement of IFT88. Each experimental run lasted ~10,000 frames before significant photobleaching. Cilia with detectable twitching or shifting motion were excluded from the analysis to avoid any effect on IFT88 dynamics (Supplemental Figure S1). Simulated experimental runs were carried out to assure our displacement and velocity analyses were valid under the conditions of the sampling rate and the resolution of our experiments (Supplemental...
FIGURE 1: sptPALM-based imaging method for dynamic studies of IFT proteins in primary cilia. (A) Isolated single-particle tracking after photoconversion. Each probe stayed in the "on" state for several tens of time points to create a short trajectory. (B) An epifluorescence image of IFT88-mEos4b showed a common bright punctum at the ciliary base (arrow), illustrating the high density of IFT proteins. (C) Each trajectory extracted from the defined curve along a primary cilium (blue arrow) was mapped to a trail in the kymograph (C, right); this is different from a wide-field (WF)-based kymograph, which usually cannot resolve individual molecules at the ciliary base (C, left). (D) A sectioned kymograph of thousands of IFT88-mEos4b trails was reconstructed and extracted from the line shown in the live-cell PALM image (panel v). One stationary and three moving IFT proteins (observed along the axoneme) are shown in panels i and ii–iv, respectively. Panel ii represents the retrograde motion, while the anterograde motion is shown in iii and iv. Scale bars: 1 μm (B); vertical, 1 s; horizontal, 2 μm (D, main panel); vertical, 500 ms; horizontal, 1 μm (D, i–iv) and 500 nm (D, v).

The DAPs and the TZ accommodated not only axonemal but also transverse IFT88 movement

The recording of short localized trajectories for IFT88-mEos4b molecules enabled statistical analysis of their displacement directions at the ciliary base. All localizations of IFT88-mEos4b of one experimental run were overlaid as the outline of the cilium, with three trajectories showing possible paths of IFT88 at the ciliary base illustrated (Figure 2A and Supplemental Movies S2–S5). Despite dynamic transport of IFT88 proteins, the image with overlaid trajectories revealed the envelope of IFT88 axonemal distribution within the CC and the three-puncta organization at the ciliary base, agreeing with our previous findings (Yang et al., 2013, 2015). One of these example trajectories showed an IFT88 molecule moving in the transverse direction from the periphery to the center of the ciliary base (Figure 2A and Supplemental Movie S3), illustrating the possibility of IFT protein movement not following the DAP or axonemal structures.

Other trajectories might not show clear direction of movement because of the combined effects of short durations and stochastic motion, as well as the noise-related uncertainty in localizing single molecules (Figure 2A and Supplemental Movies S4 and S5). Thus, to pinpoint the trajectories that represent directional movement, an effective displacement was defined by a simple linear regression fit of each trajectory of the mobile IFT proteins that spanned >45 nm (Supplemental Figure S5). Overlaid vectors of effective displacements from multiple primary cilia (straightening and alignment procedure shown in Supplemental Figure S6) showed surprising orientation differences between the cilium and the base. Most mobile IFT88 molecules within the CC moved approximately parallel to the longitudinal direction, while mobile IFT88 at the base, including the TZ and the DAPs, traveled at various angles (Figure 2, B and C, and Supplemental Figure S7). Within the DAPs and TZ, we observed...
FIGURE 2: sptPALM trajectory analysis of IFT88 at the base of primary cilia showing distinct patterns of speed and direction of motion. (A) Live-cell localizations and three representative trajectories of IFT88-mEos4b. The overlaid image shows the envelope of the IFT88 axonemal distribution (left). Three trajectories of IFT88 at the ciliary base (right) show the paths of stationary and mobile IFT proteins. (B) Overlaid vectors representing effective displacements from one cilium revealed distinct angular distributions between cilium and base. The cilium (CC) had a large population of small-angle vectors, whereas the base exhibited a diverse angular distribution. (C) Detailed angular analysis from multiple primary cilia (2786 trajectories, 18 cilia) showed that the percentage of small-angle displacements (\(< 30°\)) increased in the proximal–distal direction: DAPs < TZ < CC. The numbers of trajectories are 532, 563, and 1809 for DAP, TZ, and CC, respectively. (D) The effective velocity field calculated from the effective displacement and total travel time. (E) Representative fast-moving IFT proteins observed in the CC. (F) Statistical analysis of the orientation dependence of the effective velocity at the base were performed to compare three angular populations based on 2786 trajectories. The result suggests that high-speed IFT movement occurred when moving approximately parallel to the axoneme. The numbers of trajectories are 1333, 829, and 623 for 0–30, 30–60, and 60–90 degrees, respectively. Statistics notation: ****, \(p < 0.0001\); **, \(p < 0.01\); ns, not significant. (G) Cumulative population plot of lateral speed (Vx) shows that it was higher in the DAP and TZ regions than in the CC. The numbers of trajectories are 517, 545, and 1713 for DAP, TZ, and CC, respectively. (H) The upper bound of the Vx distribution decreased with longitudinal position. (I) Axial view of sptPALM-based analysis revealed that IFT88 proteins of nonciliated cells could travel radially between the DAPs and axoneme as well as moving tangentially. Scale bars: 500 nm (A); 100 nm (A, enlarged panel, D); 200 nm (B).

IFT proteins had a lower transverse velocity toward the ciliary compartment

The effective velocity calculated from the effective displacement indicated that some IFT proteins moved faster than others in a location- and orientation-dependent manner (Figure 2D). Fast-traveling IFT proteins in the CC had a mean effective velocity of ~600 nm/s (Figure 2E), consistent with the velocity reported in the literature (Tran et al., 2008; Besschetnova et al., 2010), confirming that our assay did not perturb IFT dynamics. Note that imaging of this work based on two-dimensional (2D) projection of three-dimensional (3D) data might potentially hide a certain fraction of information of the motion in the z direction. This may cause an underestimation of transport rate of molecules by a factor of \(\sqrt{2}/3\) on average for an axially symmetric distribution of movement, because only the speed as projected onto a 2D plane was measured. However, because of the approximately cylindrical symmetry of a cilium, it is unlikely that a large proportion of molecules traveling orthogonal to the imaging plane. Therefore, although the quantitative measurements may not be precise, the qualitative conclusions of results here and following were not affected by the 2D projection of 3D data. Analyzing the orientation dependence of the effective velocity at the base, we found that high-speed IFT movement (e.g., \(V > 500\) nm/s) occurred statistically more frequently when motion was approximately parallel to the longitudinal axis of the cilium (0/30°; Figure 2F), possibly suggesting that axoneme-associated translocation exhibited high speed, but it happened only occasionally. The fast-moving IFT88 at the base was slower than its fast-moving equivalents in the CC, and this may be due to the more crowded and intricate architectures of the DAP and TZ hindering movement.

To investigate the dependence of transverse velocity on longitudinal position, we extracted the lateral component of velocity (Vx) for each moving trajectory. Consistent with our findings regarding displacement, IFT88 exhibited higher lateral speed in the DAP and TZ regions than in the CC (Figure 2G). Interestingly, IFT proteins still...
possessed a considerable lateral speed at the TZ, suggesting that the space close to the Y-links allows lateral movement (Figure 2H). Overall, our sptPALM-based study revealed distinct patterns for IFT88 transport along primary cilia, implying different molecular hindrances due to varying molecular architectures or population densities need to be tested at different locations.

A localized axial view revealed radial transport of IFT proteins at the DAPs

Although the lateral view showed the transverse movement of IFT proteins at the DAP region, it was unclear whether these proteins traveled along the periphery of the DAPs or radially between the DAPs and the axoneme/centriole. An axial view of IFT88 movement (here molecules were observed in nonciliated centrioles) revealed that some IFT proteins moved tangentially to the periphery, while others moved radially. Therefore, IFT proteins could indeed travel between the DAPs and the axoneme through the space in between (Figure 2I), and the directions of travel spanned all angles, suggesting that the movement was not guided by the pinwheel-like DAP blades. We also used fixed-cell superresolution microscopy to show that the distribution patterns of IFT88 close to the DAPs were similar for both ciliated and nonciliated centrioles (Supplemental Figure S9), suggesting IFT88 traveled in random orientations in this region regardless of whether the cilium formed or not. This observation is consistent with our previous finding that the DAP matrix is where IFT88 proteins settle (Yang et al., 2018), implying the matrix is a region that allows dynamic movement of proteins within.

Diffusion analysis and consecutive displacement analysis indicated that IFT protein movement was more diffusive at the proximal TZ than at the distal TZ

To further refine the spatial resolution of our dynamic characterization, we performed frame-by-frame consecutive displacement analysis for each trajectory. The map of the displacement of every two consecutive points from a trajectory showed location-dependent displacement of IFT proteins (Figure 3A). A relatively large fraction of large displacements were detected at the proximal region of the TZ (Figure 3B), suggesting that the proximal TZ may support a fast-moving environment for IFT proteins, significantly different from the distal TZ region, where smaller translocation steps were observed (Figure 3C).

**FIGURE 3:** Consecutive displacement and diffusion analyses showing longitudinal dependence of IFT protein dynamics. (A) Consecutive displacement map of IFT88 from data of three cilia displayed differential characteristics of movement between the proximal TZ (pTZ) and the distal TZ (dTZ). (B) Localization map (blue) and consecutive displacements (pink) from data in A, revealing that small consecutive displacement (<45 nm) was relatively populated at the DAP and the dTZ (middle). The numbers of steps calculated are 1085, 235, 637, and 3619 for the DAP, pTZ, dTZ, and CC, respectively. (C) Statistics of consecutive displacement calculated from ~18,000 translocation steps of ~2800 trajectories in 18 cilia showing different translocation step sizes of the DAP, pTZ, dTZ, and CC. The numbers of steps calculated are 7185, 4003, 3675, and 3618 for the DAP, pTZ, dTZ, and CC, respectively. (D) Relationship between mean local displacement and localization density illustrating that the high population regions accommodate slow-moving molecules. Solid line represents the fitting result. (E) Classification of each trajectory based on diffusion characteristics ($\alpha$) was mapped onto an averaged superresolved image of a primary cilium to show its location dependence. (F) Cumulative probability distribution plot of $\alpha$ value from the data based on 15 cilia (663 trajectories) and (G) statistical analysis of longitudinal position dependence showed that IFT protein trafficking was less diffusive at the dTZ and DAP but more diffusive at the pTZ and CC. The numbers of trajectories are 255, 128, 136, and 144 for the DAP, pTZ, dTZ, and CC, respectively. Scale bar: 100 nm (A). Statistics notation: ****, $p < 0.0001$; **, $p < 0.01$, Student’s t test. Error bars represent SD.
The distal TZ, which is close to the location of the Y-links, might contain molecules hindering the motion of IFT proteins. Relatively small displacement was also observed at the DAP region (Figure 3, B and C). Analyzing location-dependent consecutive displacements, we found that regions with a higher IFT88 population density accommodated molecules that have a smaller mean consecutive displacement, and vice versa (Figure 3D), supporting the hypothesis of obstructing movement at the distal TZ and the DAP region.

Diffusion analysis based on mean square displacement (MSD; see Materials and Methods) further supports our observation in consecutive displacement analysis, revealing distinct diffusion mechanisms at the ciliary base (Figure 3, E and F). We found that IFT proteins moved more directionally in the CC, whereas diffusion modes were considerably more heterogeneous in the TZ and DAPs. Statistical analysis of the dependence of the diffusive parameter $\alpha$ on longitudinal position showed that IFT protein movement was less diffusive at the distal TZ than in the neighboring regions (Figure 3G).

Combining the comparison of consecutive displacement and diffusion parameter, we thus concluded that IFT88 molecules change dynamic characteristics multiple times along the ciliary path, moving slowly in the DAP region, relatively fast in the proximal TZ, slowly again in the distal TZ, and then much faster in the CC.

Two-color superresolution imaging revealed that the distal TZ is a potential assembly site

To understand why DAPs, proximal TZ, and distal TZ had different IFT88 dynamic characteristics, we examined where IFT88 might interact with other proteins, specifically KIF3A, the kinesin responsible for anterograde movement carrying IFT proteins in the CC, and BBS2, one of the BBSome components serving as an adaptor between IFT88 and cargo proteins (Nachury et al., 2007), in fixed RPE-1 cells. Two-color DSTORM superresolution imaging of BBS2 and IFT88 revealed that BBS2 molecules were highly populated in the CC and distal TZ, but they were considerably sparse in the proximal TZ and absent in the DAPs (Figure 4A). An image overlaying...
data from multiple cilia clearly illustrates differential localization of IFT88 and BBS2, with overlapping region of these two proteins primarily at the distal TZ, and some in the CC (Figure 4B). That is, the distal TZ, or possibly the CC, could serve as an assembly site for IFT88 and BBS2. This result might explain why we saw the slowest motion of IFT88 in the distal TZ.

Two-color superresolution images of Kif3A and IFT88 showed that, although we could see Kif3A molecules distributed along the cilia as expected, the population of KIF3A was considerably sparse in the region where IFT88 molecules were highly populated (Figure 4C). That is, only a finite number of molecular motors were available to carry IFT88 molecules into the CC. Population analysis of IFT88, BBS2, and KIF3A along the longitudinal axis shows relative populations that might reflect the assembly probability in different locations (Figure 4D). As KIF3A was less populated in the ciliary base than IFT88, KIF3A should be the limiting factor for their assembly. The ratio of KIF3A/IFT88 implies that the relative probability for each IFT88 molecule to find a KIF3A molecular motor was higher toward the distal TZ.

**DISCUSSION**

Ciliogenesis requires IFT within the CC, and this in turn requires IFT protein recruitment to the ciliary base, followed by IFT complex assembly, IFT protein–motor assembly, track engagement onto the axoneme, integration with cargoes, and ciliary entrance screening. All these processes occur at the ciliary base, where ensemble-based images show a highly crowded population of IFT proteins, whose individual trajectories of motion are impossible to delineate by conventional microscopy. This makes elucidation of these processes very challenging. By tracking stochastically photoconvertible molecules that can be isolated from one another for a short time, we demonstrated that stiPALM largely meets the needs of studying IFT protein dynamics at the ciliary base. Although individual mEos4b-IFT88 trajectories were relatively short, analysis of the aggregated trace data was sufficient to reveal the characteristics of spatiotemporal dynamics of IFT proteins at the base of mammalian primary cilia.

To better understand our results on the ciliary structural framework, we mapped the localizations onto the superresolved molecular architecture of the TZ and DAPs to find location-dependent dynamic characteristics of IFT proteins on the ciliary structural framework (Figure 4, E–G; Yang et al., 2015). Mobile IFT88 proteins travel slowly at the DAP region, fast at the Cep290-neighboring proximal TZ, very slowly through the distal TZ close to the Y-links, and very fast in the CC (Figure 4G).

Mobile IFT88 proteins traveled between DAPs and the axoneme according to varied patterns, moving in different directions with a relative slower speed when compared with IFT88 proteins in other regions. This means some areas between the DAP tips and the centriolar axoneme can accommodate IFT88 movement in constrained dynamic characteristics, potentially suggesting that some areas in the DAP region may allow IFT88 movement in an obstructive environment. The obstruction may come from the pinwheel-like structure of the DAPs or some other densely distributed molecules not apparent in the electron micrographs. This finding is not only consistent with our finding that the DAP matrix acts as the transmembrane protein gate (Yang et al., 2018), potentially pausing IFT88 movement through screening, but also agrees with the finding of a second periciliary diffusion barrier of membrane proteins at this level (Ye et al., 2018).

IFT88 in the proximal TZ, where Cep290 is localized, exhibited dynamics that were considerably different from those in adjacent regions. As IFT proteins entered the proximal TZ, they moved faster relative to the adjacent DAP and distal TZ regions in a less constrained manner. They were also less concentrated in this region than in the DAP region and the distal TZ. We hypothesize that the proximal TZ has fewer obstructive structures or it is not an assembly site for protein complexes, so IFT proteins do not get held up in this region. This is the location Ye et al. (2018) proposed as the intermediate compartment. Widely distributed IFT88 proteins, filling the space between the axoneme and the ciliary pocket, show that they are not necessarily engaged onto microtubule tracks at the proximal TZ, suggesting that a proportion of the IFT proteins in this region may not integrate with motor proteins.

The distal TZ is another region where mobile IFT proteins slow down. This is the area where several TZ proteins such as TMEM67, TCTN2, MKS1, and RPGRIP1L are found and Y-links are located (Yang et al., 2015), suggesting that IFT proteins may dwell in the region close to the Y-links. That is, this region constrains their motion to a greater degree, possibly owing to the structural hindrance provided by Y-links. That IFT proteins are held up in this area suggests that some functional activities may occur here, possibly assembly of IFT proteins with other molecules, such as molecular motors or cargoes, or entrance screening performed by Y-link proteins (Reiter et al., 2012). Several dynamic studies have found the TZ to be the diffusion barrier (Nachury et al., 2010; Chih et al., 2011; Kee et al., 2012; Ye et al., 2013, 2018; Oswald et al., 2018), and our study further confines the region to the distal TZ. Once IFT proteins pass through the TZ and enter the CC, their motion becomes faster and more unidirectional, suggesting full integration with molecular motors engaged onto the microtubule tracks. Our angular distribution analysis for each region shows that more and more IFT proteins move in the longitudinal direction as one goes from the DAPs to the TZ to the CC, suggesting that motor assembly and axonemal engagement increases in the proximal–distal direction.

Our studies allow us to reject some hypotheses and find support for others. One surprising finding is that IFT88 traveling from the recruiting site to the axoneme does not follow the “blades” of DAPs (Figure 4H). Instead, IFT88 molecules move in random directions when observed in axial view, potentially migrating with the DAP matrix region (Yang et al., 2018). Another finding is that IFT88 molecules can wander in the space enveloped by the DAPs, the axoneme, the ciliary membrane, and the Y-links. Unexpectedly, they are not entirely axoneme bound or membrane bound when traveling in the proximal TZ region, but are moving stochastically within this volume. Finally, for complex assembly, it has been shown that various BBS proteins, including BBS2, travel processively in the CC (Wei et al., 2012). We found that the assembly of IFT88 and BBS2 most likely occurs at the distal TZ region (Figure 4I). It is nearly impossible for this assembly to happen at the DAPs, because BBS2 is absent there. Based on the results presented here, we propose a model summarizing different dynamic characteristics of IFT88 at the DAPs, the proximal TZ, the distal TZ, and the CC (Figure 4J).

**MATERIALS AND METHODS**

**Cell lines and culture**

Human retinal pigment epithelial cells (hTERT-RPE-1) were cultured in DMEM/F-12 (11330-032; Thermo Fisher Scientific [Life Technologies]), Waltham, MA) supplemented with 10% fetal bovine serum (FBS) (10437-028; Thermo Fisher Scientific) and 1% penicillin–streptomycin (15140-122; Thermo Fisher Scientific [Life Technologies]). HEK293T cells were cultured with DMEM (11965-084; Thermo Fisher Scientific [Life Technologies]) with 10% FBS and 1% penicillin–streptomycin. All cells were cultured at 37°C under 5% CO₂.
Cloning and lentivirus generation
IFT88-mEos4b plasmids were generated by replacing the green fluorescent protein (GFP) sequence in the pLVX-tight puro human IFT88-GFP plasmid (Tet-On), a gift from W. J. Wang (National Yang Ming University, Taipei, Taiwan), with the mEos4b sequences from the pRSETa_mEos4b plasmid (S1073; Addgene, Cambridge, MA). Briefly, the insert (mEos4b) was amplified with a forward primer containing an MluI restriction site and a reverse primer containing a BsrGI restriction site. The primer sequences were 5′-ACGGTGCCACCATGGTGAGTTGCAGT-3′ (forward) and 5′-GGCTGTTACATTCATCGTCCGACTTGCAGG-3′ (reverse). Both the insert and the vector (IFT88-GFP) were digested with MluI (R0198; New England Biolabs, Ipswich, MA) and BsrGI (R0575; New England Biolabs) before ligation. The ligated product was transfected into DH5α-competent cells (RH617; RBBC Bioscience, Chungho, Taipei, Taiwan) to obtain the desired clone. The plasmid sequence was confirmed using DNA sequencing. An IFT88-mEos4b lentivirus was generated by cotransfection of the IFT88-mEos4b plasmid with packaging and envelope vectors (PT-5144-1; Clontech, Mountain View, CA) into HEK293T cells using a TransIT-T293 Transfection Reagent Kit (MIR 2700; Mirus, Madison, WI). The cell culture medium containing the virus was collected 3 and 4 d after transfection. The medium was centrifuged and filtered to remove cell debris. The virus was kept at −80°C until later use.

Stable clone generation and sample preparation
For generation of RPE-1 cells stably expressing IFT88-mEos4b, cells were infected with the IFT88-mEos4b lentivirus and then selected with 5 μg/ml puromycin (p8833; Sigma-Aldrich, St. Louis, MO) for 2 wk to obtain a stable cell pool. Medium was refreshed every 3 d. For creation of a single clone, the cell pool was distributed into the wells of a 96-well plate at a density of 1 cell per well and selected with puromycin for 2 wk. For preparation of a sample for localization study, cells were seeded onto poly-l-lysine (P1274; Sigma-Aldrich)-coated coverslips, and were starved for 2 d before fixation to promote cilia formation. For sptPALM imaging, samples were incubated with the addition of doxycycline (D1822; Sigma-Aldrich) to induce IFT88-mEos4b expression and then loaded into an imaging chamber (CM-B18-1; Live Cell Instrument, Nowon-gu, Seoul, Korea) filled with serum-free medium. For testing the blinking efficiency of mEos4b, cells were fixed with ice-cold methanol (32213; Sigma-Aldrich) at −20°C for 5 min before imaging.

Immunofluorescence
Cells were fixed with 4% paraformaldehyde at room temperature or methanol at −20°C for 10 min after quick rinses with phosphate-buffered saline (PBS). The cells were then permeabilized with washing solution (PBS with 0.1% Triton X-100) for 10 min and blocked with 3% bovine serum albumin for 30 min. Primary antibodies (goat IFT88 [1:100] [ab42497; Abcam, Cambridge, UK]; rabbit KIF3A [1:200] [ab11259; Abcam]; and rabbit BBS2 [1:100] [11188-2-AP; Proteintech, Rosemont, IL]) were diluted in the blocking solution. The cells were incubated in the primary antibody solution for 1 h at room temperature; this was followed by three rinses in the washing solution. The cells were then stained with secondary antibodies Alexa Fluor 647 (A21447; Thermo Fisher Scientific) and Cy3B (conjugating Cy3B maleimide; PA63131; GE, Pittsburgh, PA) to immunoglobulin G antibodies (711-005-152; Jackson, West Grove, PA) for 1 h before being washed five times.

Single molecule–localization imaging
sptPALM and dSTORM imaging were performed on a modified inverted microscope (Nikon Eclipse Ti-E; Nikon, Tokyo, Japan). The excitation lasers (OBI 637 LX 140 mW and OBI 488 LX 150 mW; Coherent, Santa Clara, CA; and Jive 561 150 mW; Cobolt, Solna, Sweden) and the activation laser (OBI 405 LX 100 mW; Coherent) were coupled into a single-mode fiber and focused on the back aperture of a high numerical aperture objective (100x/1.49 NA Apo TIRF; Nikon) for wide-field illumination. Samples were placed on a motorized three-axis stage incorporated with a Perfect Focus System (PFS; Nikon) to maintain a drift-free focus. For sptPALM imaging, the cells containing mEos4b-IFT88 were first imaged with a 488-nm laser to search for primary cilia and then illuminated with an intense (1–2 kW/cm²) 561-nm laser. For dual-color dSTORM imaging, Alexa 647 and Cy3B were imaged sequentially with 637- and 561-nm light (2–5 kW/cm²), respectively. A weak 405-nm light (<5 W/cm²) was used to activate a portion of the probes converted from a dark state to an excitation state. During the acquisition, fixed samples were incubated in an imaging buffer containing Tris-sodium chloride buffer at pH 8.0 and an oxygen-scavenging system (70– 80 mM mercaptoethanol, 0.5 mg/ml glucose oxidase, 40 μg/ml catalase, and 10% [wt/vol] glucose) (Sigma-Aldrich). The fluorescence signals were cleaned by emission filters (525/50, 593/40, and 700/75 nm; Chroma, Bellows Falls, VT) and imaged using an EMCCD (Evolve 512 Delta; Photometrics, Tucson, AZ) with a pixel size of 93 nm. For sptPALM, 5000–10,000 frames were recorded at an acquisition rate of 33/s; and for dSTORM, 10,000–20,000 frames were taken at a rate of 50/s. The peaks of individual single molecules were localized using the MetaMorph superresolution module (Molecular Devices, Sunnyvale, CA). During the live-cell imaging, RPE-1 cells were maintained in DMEM/F-12 without phenol red and incubated in a chamber heated at 37°C (TC-A/1CU-S01; Live Cell Instrument). Each sample was imaged for no more than 1 h. To systematically minimize the influence of temperature-induced drift, we added fluorescent fiducial marker (Tetraspeck; Thermo Fisher) onto both live and fixed samples at 1/200 dilution to detect drift during acquisition for postacquisition correction with ImageJ.

Analytical methods
Single-molecule tracking. Single-molecule tracking was performed via the MetaMorph superresolution module based on wavelet segmentation (Izeddin et al., 2012; Kechkar et al., 2013) and simulated annealing algorithms (Racine et al., 2006; Sibarita, 2014) for particle localization and tracking (running with a minimum length of 5 frames and a maximum search distance of 2 pixels from current point to next point) to obtain trajectories. The data from quasi-stationary cilia during an ~5-min acquisition were selected by evaluating their accumulative deviation from the nominal width and their localization pattern. A systematic characterization of the shifting effect under the quasi-stationary condition is shown in Supplemental Figure S1. In general, the shifting timescale is two orders of magnitude slower than that of active transport of IFT molecules.

Effective displacement and effective velocity analysis. Every trajectory was linearly fitted to determine its initial and final location and then described as the effective displacement (Supplemental Figure S5). For generating the effective displacement plot, as shown in Figure 2B or Supplemental Figure S7, data from different primary cilia were combined into the same coordinate system. To do this, a coordinate system adapted according to the orientation of a cilium was defined at its base. Each of the primary cilia was fitted with a spline curve and then straightened according to this curve in ImageJ. Images were aligned vertically and laterally with respect to the widest axial position at the DAPs and the spline curve. All trajectories containing the location information of...
particles were transformed onto the new coordinate system (Supplemental Figure S6). To ensure a meaningful quantification of displacement for each trajectory, we only picked trajectories with effective displacement greater than 45 nm, that is, the system resolution. The effective velocity is calculated based on the effective displacement, defined as effective displacement over duration of the corresponding single-molecule emission.

**Kymograph analysis.** For kymograph analysis shown in Figure 1D and Supplemental Figure S4, single-molecule signals were extracted only along the fitted curve for a given cilium where the spline curve starts from the base of the cilium. Kymographs were then generated from video streams of 5000–10,000 frames via Multi Kymograph (ImageJ) with a line width of 3 pixels to identify single-molecule events. To create kymographs like the one in Figure 1D, a long kymograph was reshaped into several segments combined. Given the direction of a spline curve, trails of a positive slope indicate the movement from the cilium tip to the base and vice versa. A vertical trail indicates the stationary movement along the defined curve.

**Consecutive displacement analysis.** For the consecutive displacement (translocation) analysis shown in Figure 3A, the displacements were calculated based on each distance between two consecutive locations of a trajectory that was generated as described earlier. Thousands of consecutive displacements from multiple cilia were transformed to the same coordinate before being combined. To create a map of consecutive displacements as shown in Figure 3A, we used the middle points of individual displacements as their presented locations and generated a scatter plot from these locations and the corresponding displacements. For the statistical analysis, we extracted the data from four different ciliary compartments based on the pattern of IFT88 (Yang et al., 2015); that is, \( y = [-100 \text{ nm}, 100 \text{ nm}] \) for DAP; \( y = [100 \text{ nm}, 200 \text{ nm}] \) for pTZ; \( y = [200 \text{ nm}, 400 \text{ nm}] \) for dTZ; \( y = [400 \text{ nm}, 2000 \text{ nm}] \) for CC.

**MSD and diffusive parameter \( \alpha \) analysis.** To gain insight into diffusive characteristics of molecule transport, we further computed the MSD of the IFT protein for each trajectory according to the following formula (Meijering et al., 2012):

\[
\text{MSD}(n) = \frac{1}{N-n} \sum_{i=1}^{N-n} d^2(P_i, P_{i+n})
\]

where \( N \) is the total number of time points of a trajectory; \( n \) is the number of time intervals; and \( d(P_i, P_{i+n}) \) is the distance between two points \( P_i \) and \( P_{i+n} \).

MSD is a function of time lag and allows one to characterize the mode of motion. We adopted a general formula for describing the relationship between the MSD, the diffusion parameter (\( \alpha \)), and the diffusion coefficient (\( D \)) given below for 2D particle tracking (Meijering et al., 2012):

\[
\text{MSD}(t) = 4Dt^\alpha
\]

where \( t \) is the time lag and the diffusion mode is dependent on \( \alpha \), with \( \alpha < 1 \) characterized as subdiffusion and \( \alpha > 1 \) as superdiffusion. The diffusion analysis herein enables the classification of IFT particle movement into superdiffusion, movement that behaves like directional motion driven by active transport, and subdiffusion, movement that behaves like spatially confined motion, possibly due to obstruction. Thus, we first plotted the MSD curve \( t \) for each trajectory and then fitted the curve with the above equation to calculate the \( \alpha \) value. For the optimal estimation of diffusion mode, we calculated the optimal fitting point for each MSD curve based on an empirical approximation (Michalet, 2010). The data in Figure 3, E–G, were analyzed by screening trajectories with a length of more than 7 frames and an \( \alpha \) value of 0–2.5 to minimize fitting errors and artifacts.

**Assay for the effect of pH on single-molecule event.** We examined the blinking characteristics of mEos4b in fixed samples at four different pH conditions (6, 7.2, 8, 9) in a PBS-based buffer. Each sample was imaged for 5000–10,000 frames, and then the acquired stream video was analyzed to generate a kymograph (Supplemental Figure S3A). For determining whether each blinking signal is spatially separate, a raw kymograph was first filtered to smooth the discrete signals caused by fast blinking within each trail. The image was then converted to a binary format for region size analysis and for each trail yielded a width and a length for which the width indicated the degree of signal overlap and the length was proportional to the duration of an emitter (Supplemental Figure S3D). A threshold of 6 pixels in width was chosen to differentiate between the defined single-emitter events and multi-emitter events. The ratio at different pH levels is shown in Supplemental Figure S3B. The fraction of single-emitter events of more than 10 frames is compared in Supplemental Figure S3C.

**Statistical analysis**

Our experimental data were analyzed based on eight biologically independent experiments. Quantified values that passed the statistical assumption test were subjected to the two-tailed Student’s t test using GraphPad Prism. The \( n \) values used to derive statistical analysis are noted in the figure legends.

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