Intraflagellar transport is essential for mammalian spermiogenesis but is absent in mature sperm

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Intraflagellar transport is essential for mammalian spermiogenesis but is absent in mature sperm

Jovenal T. San Agustín, Gregory J. Pazour, and George B. Witman

INTRODUCTION

Intraflagellar transport (IFT) is essential for the assembly and maintenance of most eukaryotic cilia and flagella (reviewed in Rosenbaum and Witman, 2002; Scholey, 2003). During IFT, large particles are transported along the axonemal microtubules from the cell body to the tip of the flagellum and then back to the cell body, where there is a large pool of IFT particles. The IFT particles carry cargo for assembly and maintenance of cilia and flagella (Piperno et al., 1996; Hou et al., 2007; Wren et al., 2013; Craft et al., 2015) and also carry signals between the cilium and the cell body (Pan and Snell, 2003; Goetz and Anderson, 2010; Keady et al., 2012; Liem et al., 2012; Eguether et al., 2014).

The anterograde, base-to-tip movement of IFT particles is propelled by the microtubule motor kinesin-2 (Kozminski et al., 1995; Scholey, 1996). In Caenorhabditis elegans, a second kinesin (OSM-3) cooperates with heterotrimeric kinesin-2 in this process (Ou et al., 2005). The movement of particles back to the base of the axoneme is driven by cytoplasmic dynein 2, also known as dynein 1b in vertebrates (Pazour et al., 1999; Porter et al., 1999). The IFT particles are composed of at least 16 polypeptides in IFT complex A (IFT-A; Cole et al., 2010) and also carry signals between the cilia and the cell body (Pan and Snell, 2003; Goetz and Anderson, 2010; Keady et al., 2012; Liem et al., 2012; Eguether et al., 2014).

In vertebrates, IFT also is required for assembly of motile cilia in Chlamydomonas (Kozminski et al., 1995; Pazour et al., 1998, 1999, 2000; Brazelton et al., 2001), Tetrahymena (Brown et al., 1999; Murcia et al., 2000; Pazour et al., 1998, 1999, 2000; Brazelton et al., 2001), Trypanosoma (Davidge et al., 2006; Absalon et al., 2008), and sea urchin (Morris and Scholey, 1997) and for assembly of dendritic cilia in C. elegans (Collet et al., 1998; Signor et al., 1999; Wicks et al., 2000) and Drosophila melanogaster (Han et al., 2003; Sarpal et al., 2003). In vertebrates, IFT also is required for assembly of motile cilium (Banizs et al., 2005; Kramer-Zucker et al., 2005; Gilley et al., 2014) and neoronal cilium (Davenport et al., 2007), along with photoreceptor outer segments and other sensory cilia (Marszalek et al., 2000; Pazour et al., 2002; Tsujikawa and Malicki, 2004), of nodal and other embryonic cilium (Nonaka et al., 1998; Marszalek et al., 1999; Murcia et al., 2000; Bangs et al., 2015), and of kidney, pancreatic, and skin primary

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cilia (Pazour et al., 2000; Cano et al., 2004; Lehman et al., 2009). IFT also is necessary for the maintenance of flagella in Chlamydomonas (Kozminski et al., 1993; Engel et al., 2012; Witman, 2012).

In contrast to the importance of IFT for the assembly of cilia in most somatic tissues, Plasmodium falciparum has flagellated gametes, yet lacks IFT genes. In this species, it is believed that axonemal assembly occurs in the cytoplasm and the axoneme does not become membrane enclosed until after assembly (Avidor-Reiss et al., 2004). Although IFT is needed for assembly of sensory cilia in D. melanogaster, it is dispensable for sperm flagellar assembly in this organism also. In Drosophila, it is believed that the sperm axoneme is assembled in the cytoplasm and, as in Plasmodium, does not become membrane enclosed until after assembly (Han et al., 2003; Sarpal et al., 2003). The observation that germ cell flagellar assembly does not require IFT in at least two species raises the question of whether IFT is necessary for the assembly and maintenance of the mammalian sperm flagellum. A study found that depletion of Kif3A, a subunit of kinesin-2, caused abnormalities in sperm head and tail development (Lehti et al., 2013); however, Kif3A is involved in a number of cellular processes, including membrane organelle transport and microtubule dynamics (Yamazaki et al., 1995; Boehlke et al., 2013), so these effects cannot be attributed specifically to defective IFT.

In the mouse, complete knockout of an IFT motor or particle protein causes embryonic lethality (Nonaka et al., 1993; Engel et al., 1999). However, the Oak Ridge Polycystic Kidney (orpk) insertion mutation of the Ift88 gene (Moyer et al., 1994), which encodes the IFT-B protein IFT88 (Murcia et al., 2000; Pazour et al., 2000), is hypomorphic and is reported to result in expression of what may be an alternatively spliced mRNA (Moyer et al., 1994; Taulman et al., 2001) and a reduced amount of a smaller-than-normal IFT-B protein (Taulman et al., 2001). This apparently supports sufficient residual IFT to allow the embryo to pass through critical stages in its development, so that some mice homozygous for the mutation survive to birth and even reach adulthood. Nevertheless, cilia formation in several neonatal and adult tissues is severely impaired. As a result, the Ift88 orpk mutation of the mouse, just as their homologues do in Chlamydomonas (Cole et al., 1998). To determine how the Ift88 mutation affects this complex, we compared sedimentation properties of the IFT particle from control and mutant testes (Figure 1D). In the gradient of the wild-type extract, IFT-B, represented by IFT88, IFT57, and IFT20, sedimented at ~175, as expected; IFT-A, represented by IFT140, sedimented slightly more slowly. Of importance, the particles from the mutant testes sedimented with properties similar to those from the wild-type testes. The residual IFT88 present in the mutant testes sedimented with the other IFT-B proteins, indicating that it is integrated into IFT-B. However, in the mutant, the amounts of IFT57 and IFT20 in IFT-B are in vast excess over the IFT88. This suggests that a relatively intact IFT-B particle is assembled even when there is inadequate IFT88 to be incorporated into all the particles.

**Mouse spermatogenesis and timing of IFT88 gene expression in testis**

Previously spermatogenesis was carefully examined in the rat (Leblond and Clermont, 1952; Russell et al., 1990). Although mouse spermatogenesis has not been so thoroughly characterized, Russell et al. (1990) built on the existing literature for both rat and mouse to provide a very useful description of the events during mouse spermiogenesis. To assist in interpreting the effects of the Ift88 mutation on sperm development, we have combined the diagrams of Russell et al. (1990) with our observations to order the key steps of flagellar development during mouse spermiogenesis (Figure 2). Spermatogenesis, or the development of sperm, begins with germ cells dividing and undergoing meiosis to generate spermatids. These develop into spermatozoa through the process of spermiogenesis. Spermiogenesis can be divided into 16 steps (Arabic numerals in Figure 2), which occur in synchronous waves along the seminiferous tubules of the testes. A section through a tubule will reveal germ cells at three or four stages of spermatogenesis, with one or two of these being spermiogenic. The more mature cells are organized in a band near the central lumen, and progressively less mature cells are localized in zones progressively closer to the outer surface or boundary of the tubule. Twelve distinct morphologies of the tubule (called stages and written in Roman numerals) can be distinguished by the steps of development occurring in a given tubule cross-section. For example, a section through a stage IV tubule will reveal step 15 spermatids with their flagella extending into the lumen of the tubule and

**RESULTS**

**IFT-particle proteins are lacking from epididymal sperm but are highly expressed in testis**

In a previous study using an anti-IFT88 antibody and immunofluorescence microscopy, it was reported that IFT88 was present in mature flagellated mouse sperm, where it was concentrated in the basal body region, with lesser amounts in the axoneme (Taulman et al., 2001). To determine whether this was true also for other IFT-particle proteins, we isolated sperm from the epididymis and analyzed them by Western blotting. To our surprise, we were unable to detect either IFT-A (IFT140) or IFT-B (IFT88, IFT57, and IFT20) proteins in the epididymal sperm (Figure 1A).

To confirm that IFT proteins are expressed in testes (Taulman et al., 2001), we did Northern blot analysis of RNA isolated from mouse tissues; the blots showed that tests had high IFT88 expression relative to other tissues (Figure 1B). Western blot analysis confirmed the high level of IFT88 expression in testis (Figure 1C). This suggests that IFT is important for sperm development but not needed in mature sperm.

We next investigated how the Ift88<sup>+/g377Rpw</sup> hypomorphic mutation (Moyer et al., 1994) affects testicular expression of IFT88 and other IFT-particle proteins and their assembly into IFT-A and IFT-B complexes. Compared to wild type, IFT88 protein is greatly reduced, but not completely missing, in testis of mice homozygous for the Ift88<sup>+/g377Rpw</sup> mutation (hereafter abbreviated Ift88<sup>−/−</sup>; Figure 1C). The remaining protein appears to be full length, and, in contrast to Kierszenbaum et al. (2011), we did not detect any smaller forms in mutant testis with our antibody directed against the C-terminal end of the protein. Mouse IFT-B proteins IFT88, IFT57, IFT52, and IFT20 sediment as a 175S particle in sucrose density gradients (Pazour et al., 2002), indicating that these proteins form a large complex in the mouse, just as their homologues do in Chlamydomonas (Cole et al., 1998). To determine how the Ift88 mutation affects this complex, we compared sedimentation properties of the IFT particle from control and mutant testis (Figure 1D). In the gradient of the wild-type extract, IFT-B, represented by IFT88, IFT57, and IFT20, migrated at ~175, as expected; IFT-A, represented by IFT140, sedimented slightly more slowly. Of importance, the particles from the mutant testes sedimented with properties similar to those from the wild-type testes. The residual IFT88 present in the mutant testis sedimented with the other IFT-B proteins, indicating that it is integrated into IFT-B. However, in the mutant, the amounts of IFT57 and IFT20 in IFT-B are in vast excess over the amount of IFT88. This suggests that a relatively intact IFT-B particle is assembled even when there is inadequate IFT88 to be incorporated into all the particles.

**IFT88 in sperm development**
FIGURE 1: IFT88 expression in wild-type and Ift88−/− testes. (A) Western blots of wild-type testis and epididymal sperm extracts probed with antibodies to the indicated IFT-A and IFT-B proteins and to Cα2, the testis-specific isof orm of the catalytic subunit of cAMP-dependent protein kinase. IFT proteins are readily detected in testis extract but are not found in epididymal sperm. The blot was stripped and reprobed with an antibody to Cα2 to confirm that comparable amounts of sperm protein were loaded in each lane. (B) Northern blot of wild-type mouse tissues probed with cDNAs encoding IFT88, radial spoke protein 3, tektin, and actin. IFT88 mRNA is most highly expressed in testis, with lower levels of expression in kidney, liver, lung, and brain. Tektin and radial spoke protein 3, components of ciliary and flagellar axonemes, also are most highly expressed in testis. In contrast, actin is expressed in all tissues. (C) Top, Western blot of IFT88 protein in testis extracts. The anti-IFT88 antibody recognizes a single protein of ~90 kDa, which agrees well with the predicted mass of 92.9 kDa for IFT88, in homozygous Ift88−/− (−/−), wild-type (+/+), and heterozygous (+/−) mice, but the amount of protein in the mutant mouse testis is much lower than in the wild-type or heterozygous testes. Bottom, the same blot stripped and reprobed with the anti-Cα2 antibody as a loading control. (D) Western blots of fractions from sucrose density gradients loaded with testis extracts from wild-type (top) and Ift88−/− (bottom) mice; the blots were probed with antibodies to an IFT-A protein (IFT140), IFT-B proteins (IFT88, IFT57, IFT20), and Cα2. In the gradient of wild-type testis extract, IFT88, IFT57, and IFT20 cosediment at ~17S, whereas IFT140 sediments slightly more slowly. In the gradient of mutant testis extract, there is about the same amount of IFT140, IFT57, and IFT20 as in wild type, but the amount of IFT88 is greatly reduced. Nevertheless, all three IFT-B proteins cosediment at ~17S. The blots were probed for Cα2 as a control to confirm that equivalent amounts of wild-type and Ift88−/− extracts were loaded on the sucrose gradients.

step 4 spermatids located in a band between the lumen and the outer surface of the tubule. Spermiogenesis begins at step 1 with the appearance of haploid, round spermatids arising from two sequential meiotic divisions of the diplotene spermatocytes. During step 2-3, the 9 + 2 flagellar axoneme begins to elongate from a basal body located just below the plasma membrane and reaches nearly full length (Irons and Clermont, 1982). At this time, the axoneme is tightly surrounded by the flagellar membrane, which is continuous with cellular plasma membrane. Also during step 2-3, the precursors or “anlagen” of the fibrous sheath columns begin to form at the distal end of the flagellum (Sakai et al., 1986). The fibrous sheath columns and ribs subsequently are assembled in a distal-to-proximal direction in what will become the principal piece of the sperm (Irons and Clermont, 1982). Between steps 6 and 7 (Russell et al., 1990), the basal body with attached axoneme migrates inward to the nucleus and pulls the plasma membrane along with it to create an invagination of the plasma membrane that now surrounds the proximal portion of the flagellum. At step 8, the outer dense fibers begin to form at the proximal end of the axoneme. The fibers assemble along the axonemal doublet microtubules in a proximal-to-distal direction, eventually extending the entire length of the midpiece and principal piece but not reaching their full diameter until step 16. At step 9, the nucleus begins to elongate and condense. Also at this step, the annulus—a septin-based ring of dense material—starts to form and surrounds the axoneme at its base (Guan et al., 2009). At step 15, the annulus migrates distally along the axoneme to the future site of the midpiece/principal piece junction (Guan et al., 2009; Kwitny et al., 2010). The axoneme and outer dense fibers proximal to this site are now surrounded by the spermatid’s caudal cytoplasm and will become the midpiece. Beginning
FIGURE 2: Key steps in the development of the mouse sperm flagellum during spermiogenesis. Adapted from Russell et al. (1990). Boxed events were determined from our experiments unless stated otherwise. Steps 1–8 are round spermatids, before the beginning of nuclear condensation and elongation in step 9. Based on the rat (Leblond and Clermont, 1952; Russell et al., 1990), the axoneme is assembled to full length in step 2-3. Assembly of outer dense fibers (ODFs) around the axoneme begins at the proximal end of the midpiece of step 8 spermatids; the process is completed just before the mature sperm is released into the lumen of the seminiferous tubule. Fibrous sheath (FS) assembly begins at the distal end of the principal piece of step 2 rat spermatids with the formation of fibrous sheath columns along the axonemal doublet microtubules 3 and 8; a similar event is presumed to take place in step 2-3 mouse spermatids. Assembly of fibrous sheath ribs on the columns starts in step 10 spermatids and is complete in step 13 spermatids. The manchette is first seen in step 8 spermatids and disappears by step 14. The annulus slides distally down the axoneme to its final location between the distal end of the midpiece and the proximal end of the principal piece when step 15 is reached (Kwitny et al., 2010). The mitochondria then condense along the length of the midpiece.

at step 15, mitochondria in the developing midpiece migrate to the axoneme and condense around the axoneme and outer dense fibers (Russell et al., 1990). In step 16, the flagellar membrane becomes closely opposed to the mitochondrial sheath in the midpiece and the excess cytoplasm is eliminated as the residual body (O’Donnell et al., 2011). Mature sperm are then released into the lumen and exit the testis.

To determine when and where during spermatogenesis IFT88 is expressed in testis, we used immunofluorescence microscopy of sections through wild-type seminiferous tubules (Figure 3A). We found that IFT88 was first apparent in pachytene spermatocytes of stage II-III (Figure 3Aa) and was strongly labeled in pachytene cells at stage VIII (Figure 3Ab). It continued to be expressed as the spermatocytes underwent maturation, as shown by its presence in the tails of developing spermatids at both of these stages (Figure 3Aa, and b).

Because most sections of spermatogenic tubules contain spermatids in more than one stage (e.g., the stage II-III section
shown in Figure 3Aa contains spermatids in steps 2-3 and 14), it was difficult to determine conclusively whether IFT88 was localized to the tails of both early and late spermatids. To address this, we examined the distribution of IFT88 in isolated spermatids that could be identified with regard to step using the criteria of Russell et al. (1990; Figure 3B). The anti-IFT88 antibody strongly labeled the heads and developing flagella of step 2-3 through 11 spermatids (Figure 3B, a–e). At step 15, IFT88 labeling was less bright and concentrated in the cytoplasmic lobe and at the proximal end of the principal piece, suggesting that IFT winds down as the tail reaches maturity (Figure 3Bf). IFT88 labeling was not detected in cells that appeared to be either late step 16 spermatids or mature sperm that already had been released into the tubule lumen (Figure 3Bg).

To confirm that IFT88 was absent from fully formed sperm, we also examined sperm extruded from the caudal epididymides. Even using an antigen retrieval step similar to one used previously to enhance both IFT88 and tubulin labeling in the photoreceptor outer segment (Keady et al., 2011),

**FIGURE 3:** Timing and location of IFT88 expression in wild-type testis. (A) Immunofluorescence micrographs of sections through (a) stage II-III and (b) stage VIII seminiferous tubules labeled with anti-IFT88 antibody (green), anti-α-tubulin (red), and the nuclear stain TOTO-3 (blue). Arrows point out pachytene cell nuclei, in which the chromatin is condensed into multiple distinct foci; in a, IFT88 is just beginning to appear in the cytoplasm surrounding the pachytene nuclei. The bright red tubulin labeling marks Sertoli cells, through which the developing spermatids pass. es, elongated spermatids; f, flagella; L, lumen; p, pachytene; rs, round spermatids. Scale bar, 20 μm. (B) Immunofluorescence micrographs of individual isolated spermatids labeled with anti-IFT88 (green), anti-α-tubulin (red), and TOTO-3 (blue). (a) The round head and short flagellum suggests that this is an early step 2-3 spermatid. (b) The round head and longer flagellum indicates that this is a step 7 or possibly step 8 spermatid. (c) The nucleus has just begun to elongate, but the head is round and lacks a cytoplasmic lobe, indicating that this is a step 7 or possibly step 8 spermatid. (d) The more elongated shape of the head and nucleus and the distinct cytoplasmic lobe suggest that this is a step 10 spermatid. (e) The nucleus is highly condensed, and the head is more elongated than in d, suggesting that this is a step 11 spermatid. (f) The condensed nucleus, sharp narrow apex of the head, and presence of long narrow cytoplasmic lobe indicate that this is a later, probably step 15, spermatid. (g) The shape of the head (which is properly oriented to display the typical hook only in the spermatid on the right) and absence of cytoplasmic lobes suggest that these are either late step 16 spermatids that have already lost their cytoplasmic lobes or mature sperm from the tubule lumen. IFT88 labeling is present in the heads and tails of step 2-3 through 11 spermatids, limited largely to the cytoplasmic lobe in the step 15 spermatid, and no longer detected in the step 16 spermatids/mature sperm. Asterisks mark heads from other spermatogenic cells. Scale bar, 10 μm. (C) Immunofluorescence micrographs of epididymal sperm labeled with anti-acetylated α-tubulin (red), anti-IFT88 (green), and DAPI (blue). (a, b) Merged images; (a’, b’) just the green (IFT88) channel for the respective images. Tubulin labeling is readily apparent, but IFT88 is not detectable in the sperm head or tail. Although the distal part of the tail has doubled back on itself in the sperm shown in b and b’, the image shows what appears to be a cytoplasmic droplet (arrow) that serves as a positive control for the IFT88 labeling. Scale bar, 10 μm.
we extruded the contents of the cauda epididymides into phosphate-buffered saline (PBS) and examined them by light microscopy. The mean total count (heads plus tails plus heads with tails) of wild-type sperm were each mated to two wild-type females and litters scored during the fourth week. After 6 wk, males were separated from females for 3 wk.

**TABLE 1:** Male fertility.

<table>
<thead>
<tr>
<th>Sperm count</th>
<th>Wild-type males</th>
<th>Ift88−/+ males</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild type</td>
<td>19,000,000 ± 2,600,000</td>
<td>56,000 ± 31,000 (p ≤ 0.0001)</td>
</tr>
</tbody>
</table>

Sperm count included heads with tails, heads without tails, and tails without heads if present. Three mutants and six control males were harvested at >8 wk of age.

**TABLE 2:** Epididymal sperm count.

we detected no IFT88 in the epididymal sperm head or tail (Figure 3C).

**Male Ift88−/+ mice are sterile**

To test the importance of IFT88 to male fertility, five wild-type and five Ift88−/− male mice were each mated to two wild-type female mice. All 10 females mated to wild-type males produced litters during week 4 (Table 1). In contrast, although the females mated to Ift88−/− male mice were plugged, none became pregnant. All mice were then rested for 3 wk, and the females that originally had been mated to the mutant mice were mated to wild-type mice and vice versa. Again, all females mated to wild-type males produced litters, confirming that they were fertile. However, the females mated to the mutant males failed to produce litters, indicating that Ift88−/− male mice are sterile. In contrast, female Ift88−/− mice are fertile (McDermott et al., 2010).

To understand the origin of male infertility in the Ift88−/− mice, we extruded the contents of the cauda epididymides into phosphate-buffered saline (PBS) and examined them by light microscopy. The mean total count (heads plus tails plus heads with tails) for mutant mice was ~350 times lower than for wild-type animals (Table 2). The tails of wild-type sperm beat slowly, whereas those of the Ift88−/− sperm were completely immotile. Wild-type sperm had well-shaped heads and long, smoothly tapering flagella, whereas the mutant sperm were morphologically abnormal (Figure 4A). The mutant sperm heads often lacked flagella entirely, and when a flagellum was present, it was always short and misshapen.

The reduced sperm production was confirmed by periodic acid-Schiff (PAS)-hematoxylin–stained sections of wild-type and Ift88−/− testes. Mutants showed a reduced number of condensed spermatid nuclei in the epithelium and a distinct absence of flagella in the lumen of the seminiferous tubules (Figure 4B). In addition, the sperm heads were not uniformly located at the edge of the lumen of the Ift88−/− tubules as they are in controls (Figure 4B). The very low sperm count, abnormal sperm morphology, and lack of sperm motility undoubtedly account for the infertility of the Ift88−/− mice.
Expression of representative sperm flagella structural proteins is normal in Ift88<sup>−/−</sup> testes

To determine the effect of reduced IFT88 on the expression of structural proteins necessary for the development of sperm flagella, we used immunohistochemistry to evaluate production and localization of the fibrous sheath rib component ropporin (Fujita et al., 2000) and the outer dense fiber component Odf2 (Hoyer-Fender et al., 1998). In the wild-type testis, ropporin is first detected in the cytoplasm of step 4 round spermatids and begins to be incorporated into the flagella in step 9 spermatids (Supplemental Figure S1). Staining also is observed in the residual bodies that are resorbed by the Sertoli cells in stage VII seminiferous tubules. In mutant testes, the staining pattern was similar, except that few flagella were observed in the seminiferous tubules (Supplemental Figure S2). However, when flagella were present, they were stained by the anti-ropporin antibodies (Figure 4C).

Odf2 was initially seen in the cytoplasm of step 12 elongate spermatids (Supplemental Figure S3). Odf2 staining became detectable in the flagella of step 13 elongate spermatids and was most intense in the cytoplasm and flagella of step 15 spermatids. These data suggest that there is a substantial lag between the synthesis of Odf2 mRNA, which is reported to begin in step 5 spermatids and peak in step 10/11 elongating spermatids (Hoyer-Fender et al., 1998), and the appearance of Odf2 protein. Subsequently the Odf2 protein is moved into the flagella of step 16 spermatids; very little appeared to be resorbed in the residual bodies. In Ift88<sup>−/−</sup> testes, the few flagella that formed were positive for Odf2 (Figure 4D). Of interest, even though most Odf2 was not incorporated into flagella in the mutant, Odf2 label disappeared by step 15 (Supplemental Figure S4), suggesting that the excess protein was broken down. Thus, even though sperm development is greatly altered by the reduction in amount of IFT88, the timing and pattern of expression of important fibrous sheath rib and outer dense fiber components are not noticeably altered.

Axonemal growth is truncated while microtubules and accessory structure components accumulate in Ift88<sup>−/−</sup> spermatid flagella

To learn more about the morphological defects in Ift88<sup>−/−</sup> sperm, we carried out ultrastructural analysis of developing spermatids. Stage VI–VIII tubules, which contain spermatids at steps 6–8 and steps 15 and 16, are particularly informative for examining fibrous sheath and outer dense fiber assembly.

Cross-sections through the caudal cytoplasm of controls at these stages showed well-organized axonemes, which, at the later stages, were surrounded by outer dense fibers; in the developing midpiece, mitochondria were condensing around the axonemes and associated outer dense fibers (Figure 5, A and C). In contrast to the controls, most sections through the caudal cytoplasm of mutant sperm did not reveal any cross-sections of axonemes (Figure 5, B, D, and E); in these cells, the outer dense fibers formed ectopically in the midpiece of another Ift88<sup>−/−</sup> step 16 spermatid. Mitochondria (m) have condensed around several ectopically located outer dense fibers (white arrows). Scale bar, 500 nm. (F) A section through the midpiece of an Ift88<sup>−/−</sup> step 16 spermatid, showing one of the few instances in which an axoneme was observed. Even in the presence of an axoneme and its associated outer dense fibers, some outer dense fibers have assembled ectopically (white arrows) in the cytoplasm.

A microtubule (arrowhead) is associated with one or more of the outer dense fibers. Scale bar, 500 nm.
cytoplasm and often had mitochondria associated with them. Even when an axoneme was present, outer dense fibers, sometimes associated with microtubules, were formed ectopically (Figure 5F).

To better understand how the reduction in IFT affected assembly of the axoneme and its accessory structures, we examined the principal pieces of the relatively rare Ift88−/− spermatids that had a flagellum (Figure 6). In principal pieces of controls at step 6, outer dense fibers have not yet initiated, but the longitudinal columns of the fibrous sheath are formed and attached to doublets 3 and 8 (Figure 6I). At step 15, nine outer dense fibers are being assembled along the axoneme and will be completed in the next step. The fibrous sheath is completed at step 15 and consists of two longitudinal columns attached to outer doublets 3 and 8 and transverse ribs that form an arc around the axoneme, bridging the longitudinal columns (Figure 6, A and B). In principal pieces of the mutant, fibrous sheath assembly is defective from the earliest stages, and the longitudinal columns appear either to never form or never attach to the doublets (Figure 6, J and K). In some cases, the fibrous sheath rib material surrounded the outer dense fibers normally (Figure 6D), but in most cases, it condensed into large aggregates that were associated with the membrane (Figure 6, C, E, and G). Axonemes often were surrounded by supernumerary outer dense fibers (Figure 6, D and H); ectopic microtubules also were observed (Figure 6K). In some cross-sections of tails lacking an organized axoneme, large numbers of outer dense fibers were associated with singlet and doublet microtubules (Figure 6, E and F). In some Ift88−/− spermatids, the distal end of the flagellum was greatly swollen, suggesting an accumulation of unassembled axonemal precursors (Figure 6, C and G). Axonemes in the mutant flagella appeared to have a normal complement of inner and outer dynein arms and radial spokes (Figure 6, D, H, J, and K).

Nuclear reshaping of Ift88−/− spermatids is impaired

Normally, in late development, sperm nuclear chromatin condenses and the head undergoes an elongation event that forms the mature structure. This process can be seen in wild-type spermatids at step 11 (Figure 7A). Although Ift88−/− mutant cells undergo chromatin condensation and head elongation, many of the heads are abnormally shaped, as previously reported (Kierszenbaum et al., 2011; Figure 7B). At the electron microscopic level, control step 11 spermatids have partially condensed and elongated nuclei capped by the acrosome (Figure 7D). In addition, just distal to the acrosome, the nucleus is surrounded by the microtubular manchette. The anterior ends of the manchette microtubules are embedded in the nuclear ring at the posterior edge of the acrosome, and the microtubules extend posteriorly parallel to the elongating nucleus, ending in scattered electron densities in the caudal cytoplasm of the spermatid. Equivalent electron micrographs of Ift88−/− step 11 spermatids revealed abnormally shaped nuclei that are tipped by material remains associated with the distended flagellar membrane. (H) A higher magnification of the boxed region in G showing the axoneme, which lacks fibrous sheath longitudinal columns (arrowheads) and has supernumerary outer dense fibers. (I) Cross-section through a flagellum of a wild-type step 6 spermatozoon. The anlagen of the fibrous sheath longitudinal columns (black arrowheads) is visible adjacent to doublets 3 and 8. (J, K) Cross-sections through flagella of Ift88−/− probable step 6 spermatids. The anlagen of the fibrous sheath longitudinal columns are missing. The mutant flagellum in K has supernumerary singlet and doublet microtubules (white arrowheads). Scale bars, 1 μm (A, C, E, G), 100 nm (B, D, F, H–K).
The mutant testis is about normal but that the number of round spermatids is only about two-thirds of the wild-type value (Table 3). At least some of this loss is due to apoptosis. In wild-type testes, apoptosis of spermatogonia is observed in late development (tubule stages IX–XII), but apoptosis of spermatocytes is rarely observed (Dym, 1994; Figure 8A). In contrast, in Ift88−/− testes, numerous terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL)-positive spermatocytes at the late pachytene, diplotene, and metaphase stages were observed (Figure 8, B–D). Therefore loss of cells appears to begin after initiation of synthesis of IFT88 but before initiation of flagellar assembly.

Although apoptosis of meiotic cells appears to be one cause of the reduced sperm count in Ift88−/−, the reduction in sperm count is greater than can be accounted for by the loss of meiotic cells, so other mechanisms must be active. Indeed, sperm cells appear to be lost via phagocytosis by Sertoli cells. Normally at stage IX, mature wild-type sperm have been released into the lumen, and elongated condensed heads are not visible. At this stage, the next generation of spermatids is at step 9 and just beginning to undergo nuclear condensation and elongation (Figure 8E). In contrast to wild type, Ift88−/− tubules at stage IX contain numerous highly condensed nuclei in the seminiferous epithelium (Figure 8F). In many cases, these are deep in the epithelium, below the level of the step 9 spermatids (bracket, Figure 8F). It is likely that these represent later-step spermatids that were resorbed by the Sertoli cells during a preceding stage, preventing their release into the lumen.

**DISCUSSION**

**Reduced IFT88 impairs mouse sperm flagellar assembly**

The results of this study show that IFT88 is necessary for mammalian spermiogenesis and male fertility. This is in marked contrast to *Drosophila*, for which mutation of either Ift88 (Han et al., 2003) or a subunit of the IFT motor kinesin-2 (Sarpal et al., 2003) had no effect on sperm formation or male fertility. *Drosophila* sperm axoneme development occurs within the cytoplasm, and the axoneme is not fully enclosed by a flagellar membrane until it is fully elongated (Han et al., 2003). Thus it has been suggested that the axoneme is readily accessible to axonemal precursors in the cytoplasm and IFT is not needed (Han et al., 2003). However, this may not be completely correct, as the tip of the growing axoneme in *Drosophila* is tightly surrounded by an involution of the plasma membrane (Tokuyasu et al., 1972; Gottardto et al., 2013; see Figure 2B in Witman, 2003). This geometry is very similar to the early stages of *Chlamydomonas* flagellar growth, which cannot occur in the absence of IFT88 (Pazour et al., 2000). Therefore *Drosophila* may have evolved an alternative mechanism for delivering axonemal precursors to the membrane-enclosed tip of the elongating axoneme, perhaps to meet novel requirements posed by the assembly of an extremely long (1.8–2.0 mm) flagellum (Han et al., 2003). One possibility is that mRNAs encoding tubulin and other axonemal proteins are present and

<table>
<thead>
<tr>
<th>Preleptotene spermatocytes</th>
<th>Pachytene spermatocytes</th>
<th>Round spermatids</th>
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<tbody>
<tr>
<td>Wild type</td>
<td>12.5 ± 1.9</td>
<td>12.1 ± 1.7</td>
</tr>
<tr>
<td>Ift88−/−</td>
<td>11.3 ± 1.6 (ns)</td>
<td>11.5 ± 1.6 (ns)</td>
</tr>
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Numbers indicate cells per 3-µm-thick seminiferous tubule cross-sections ± SD (see Materials and Methods). Three animals per genotype. Significance is compared with wild type at the same stage.

**TABLE 3:** Developing sperm cell count in seminiferous tubule cross-sections.
failure to assemble may be because the mutant flagella do not become sufficiently long to initiate assembly or because longitudinal column proteins are not moved into the mutant flagellum. In contrast to the lack of longitudinal columns, the Ift88−/− flagella contain large accumulations of fibrous sheath rib material in their swollen tips. This accumulation strongly suggests that the rib material is actively transported to the flagellar tip. It may be carried by a non-IFT mechanism; if it is carried by IFT, it is not dependent upon wild-type levels of IFT88.

In the Ift88−/− spermatids, outer dense fibers assemble ectopically in the caudal cytoplasm when there is no midpiece axoneme to organize them. In these cases, the mitochondria condense around single or small groups of outer dense fibers, demonstrating that the outer dense fibers alone are sufficient to organize the mitochondrial sheath. In flagella of the mutant, outer dense fiber material appears to be in excess of assembled axonemal proteins, causing ectopic outer dense fibers to form on singlet or doublet microtubules where no axoneme is present. These observations indicate that outer dense fibers have a strong affinity for microtubules but that an intact axoneme is required to organize them into a ninefold array. Because outer dense fibers assemble from base to tip and are present in Ift88−/− sperm flagella, their precursors also are likely to be transported to their site of assembly by a mechanism not dependent on translated within the membrane cap surrounding the tip of the axoneme (Fabian and Brill, 2012), ensuring that the axonemal precursors are continuously available without the need for transport over great distances. In contrast, in the mouse, ribosomes are abundant in the spermatid caudal cytoplasm, including around the developing midpiece, but we never observed ribosomes in the developing flagellum distal to the annulus.

**Assembly of flagellar axoneme and accessory structures in Ift88−/− spermatids**

In *Chlamydomonas*, axonemal components, including tubulin, inner and outer arm dynein, nexin–dynein regulatory complex subunits, and radial spokes, are transported to the tip of the flagellum by IFT (Piperno et al., 1996; Hou et al., 2007; Wren et al., 2013; Craft et al., 2015). Mouse sperm flagella contain all of these components but also contain fibrous sheath and outer dense fiber accessory structures, which are aberrant in the Ift88−/− mouse. This is particularly apparent for the fibrous sheath. The fibrous sheath is composed of proteins organized into two longitudinal columns that run along the axoneme and are connected by semicircular ribs. The longitudinal columns normally assemble in a distal-to-proximal direction soon after formation of the full-length axoneme (Irons and Clermont, 1982), but they are never observed in the Ift88−/− flagellum. The failure to assemble may be because the mutant flagella do not become sufficiently long to initiate assembly or because longitudinal column proteins are not moved into the mutant flagellum. In contrast to the lack of longitudinal columns, the Ift88−/− flagella contain large accumulations of fibrous sheath rib material in their swollen tips. This accumulation strongly suggests that the rib material is actively transported to the flagellar tip. It may be carried by a non-IFT mechanism; if it is carried by IFT, it is not dependent upon wild-type levels of IFT88.

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**FIGURE 8:** Many Ift88−/− spermatocytes and spermatids are eliminated by resorption or apoptosis. (A, B) Sections of wild-type (A) and Ift88−/− (B) stage IX seminiferous tubules stained by the TUNEL procedure to reveal apoptotic and/or necrotic cells. No apoptotic cells were observed in the wild-type tubule, but numerous TUNEL-positive cells were present in the Ift88−/− tubule. The sections were counterstained with hematoxylin. (C, D) Higher-magnification images of Ift88−/− tubules in stages IX (C) and XI (D) revealed TUNEL-positive cells that appeared to have reached the pachytene (P), diplotene (Di), and metaphase (M) stages. (E) Cross-section of a wild-type stage IX seminiferous tubule. The mature spermatids have been released, so no condensed nuclei are visible in the seminiferous epithelium. The next generation of spermatids (bracket) is at step 9 and is nearing the tubule lumen. PAS-hematoxylin staining. (F) A comparable section through an Ift88−/− stage IX seminiferous tubule. Numerous elongated heads with condensed nuclei (arrowheads) are visible deep in the seminiferous epithelium, many even below the level of the step 9 spermatids (bracket) present at this stage, suggesting that the elongated spermatids have been resorbed by the Sertoli cells. PAS-hematoxylin staining. Scale, bars, 20 μm (A, B, E, F), 5 μm (C, D).
normal levels of IFT88. Lehti et al. (2013) reported that the outer dense fiber protein ODF3 was coimmunoprecipitated with KIF3A, raising the possibility that it is coupled directly to kinesin-2 for import into the developing flagellum.

The presence of many unorganized singlet and doublet microtubules in the Ift88−/− flagella indicates that tubulin also has been transported into the flagellum in considerable excess of what has assembled into 9 + 2 axonemes. Assuming that IFT-B lacking IFT88 is trafficked through flagella, this is consistent with the prediction of Bhogaraju et al. (2013) that two other IFT-B proteins, IFT81 and IFT74, together form a tubulin-binding module for transport of tubulin by IFT. The fact that the microtubules in the Ift88−/− flagellum often are not organized into an axoneme probably is due to a failure of IFT to deliver minor proteins that are essential for assembly of the axonemal superstructure.

IFT88 normally is expressed before the time when abnormalities appear in mutant germ cells

The synthesis of IFT88 protein was first detected in early pachytene spermatocytes, and the protein was highly expressed in midpachytene spermatocytes, well before flagellar assembly or nuclear reshaping is initiated after meiosis. Because there is no evidence that IFT88 participates in meiosis, it is probable that the germ cell makes IFT88 at this time so that there will be an ample pool available when flagellar growth is initiated shortly after meiosis. This timing of expression is only slightly earlier than that of the cAMP-dependent protein kinase catalytic subunit ζCα, which is a component of the axoneme (San Agustin and Witman, 2001) and is first expressed in midpachytene spermatocytes (San Agustin et al., 2000). It also is consistent with a report that synthesis of SDS-soluble sperm proteins is highest during meiosis (O’Brien and Bellve, 1980). Of interest, expression of KIF3A also was first detected in pachytene spermatocytes and subsequently increased (Lehti et al., 2013). The timing of expression of IFT88 is in contrast to that of the accessory structure proteins ropporin and Odf2, which were first detected in step 4 and 12 meiotic spermatids, respectively. These proteins are incorporated into the flagellum only after its elongation has been completed, so presumably there is no need to accumulate a pool of them before flagellar assembly.

Apoptosis was observed in Ift88−/− spermatocytes that had reached the late pachytene, diplotene, and metaphase stages. This is before flagellar assembly or nuclear reshaping, so defects in these processes cannot have been the cause of apoptosis in these cells. Because large amounts of IFT88 protein normally are made before or during these stages, it is likely that spermatocytes have a highly sensitive mechanism to detect the amount of IFT88 that has accumulated and respond by programmed cell death when the levels are insufficient for subsequent spermatogenic processes. If such a monitoring mechanism extends to sperm structural proteins, it might explain the absence of mature flagellated sperm in other cells in that it has virtually no cytoplasm, no protein synthesis, and presumably no pool of flagellar precursors, so it is highly improbable that it undergoes turnover of flagellar components. Moreover, although signals may move from the head to the tail and vice versa, these signals are likely to be mediated by ions or small molecules rather than IFT (Kirichok et al., 2006). Therefore the mature mammalian sperm lacks processes mediated by IFT in other cilia and flagella and, in the absence of these processes, has dispensed with the IFT system.

MATERIALS AND METHODS

Mouse breeding

Mice used for this work were a recombinant inbred line derived from C3H/HeJ and FvB/NJ containing the wild-type Pde6b (Rd1) allele from C57BL/6J and the Ift88β737Rpw mutation (Moyer et al., 1994). Mutant (Ift88−/−) and control (Ift88+/− or Ift88+/+) animals were derived from heterozygous-by-heterozygous crosses (Pazour et al., 2000, 2002).

For fertility testing, five control and five Ift88−/− males were each paired with two wild-type C57BL/6J females. Plug formation was checked daily during the first week, and litters were tracked for the duration of the study. After 6 wk, the females were separated from the males for 3 wk to ensure that paternity could be established. After this 3-wk period, the males were swapped so that females exposed to mutant males were paired with control males and vice versa. The second mating phase was terminated after 6 wk.

All mouse work was performed with approval of the University of Massachusetts Medical School Institutional Animal Care and Use Committee.

Antibodies and reagents

Rabbit antibodies against mouse IFT140, IFT88, IFT57, and IFT20 were made and affinity purified as described (Pazour et al., 2002, Jonassen et al., 2012). Affinity-purified rabbit anti-ropporin (Fujita et al., 2000) was a gift from Shuh Narumiya (Department of Pharmacology, Kyoto University Faculty of Medicine, Kyoto, Japan). Affinity-purified rabbit anti-Odf2 (Hoyer-Fender et al., 1995) was a gift.
from Sigrid Hoyer-Fender (Department of Zoology-Developmental Biology, University of Göttingen, Göttingen, Germany). The affinity-purified anti-Cαt antibody was described previously (San Agustin et al., 2000). Monoclonal anti-α-tubulin antibody (clone B-5-1-2) and monoclonal anti-acetylated α-tubulin antibody 6-11B-1 were purchased from Sigma-Aldrich (St. Louis, MO).

**Northern and Western blotting**

Northern blotting was performed according to standard procedures (Sambrook et al., 1989) using a Mouse Multiple Tissue Northern Blot (7762-1, BD Biosciences, Palo Alto, CA) and random-primed, 12P-labeled cDNA as a probe.Tests extracts and Western blots were prepared as described previously (San Agustin et al., 2000; San Agustin and Witman, 2001). Some used Western blots were stripped by overnight incubation at room temperature with Restore Western Blot Stripping Buffer (Pierce Biotechnology, Rockford, IL), treated with blocker and then reprobed with a different antibody.

**Sucrose gradient fractionation of testis extracts**

Two testes were excised, placed in 500 μl of ice-cold extraction buffer (35 mM NaHCO3, 2 mM Na2HPO4, 70 mM KCl, 74 mM NaCl [McGrady, 1979], 1 mM dithiothreitol, 0.16% digitonin, 10 mM 4-aminobenzamidine, 2 mM EDTA, 50 μM leupeptin, 15 μM pepstatin, 200 μM tosyl phenylalanyl chloromethyl ketone, 100 μM tosyl lysine chloromethyl ketone, 1 μg/ml aprotinin, final pH 7.4) and homogenized in a Potter-Elvehjem tissue grinder. The homogenate was centrifuged at 6500 × g for 15 min at 4°C and further clarified at 56,000 × g for 1 h at 4°C. The clarified testis extract was layered on a 5–20% sucrose gradient that had been preformed in a Beckman L5-75 ultracentrifuge (36,000 rpm, SW41 rotor). The cumulative centrifugal effect (a2t) for all runs was set at 6.4 × 1010 rad²/s. Sedimentation standards were thyroglobulin (19.4S), catalase (11.3S), and bovine serum albumin (BSA; 4.4S; Cole et al., 1998). Sucrose gradient fractions (500 μl) were collected, and the proteins were separated by SDS-PAGE, transferred to polyvinylidene fluoride membranes, and probed with anti-IFT antibodies.

**Examination and counting of epididymal sperm**

Cauda epididymides were excised and cleared of fatty tissue. The vas deferens was cut, with a short portion left to serve as outlet for the sperm. The sperm were then prodded out of the epididymis with a blunt 21-gauge needle into 1 ml of PBS and counted using a hemocytometer. For morphological examination by differential interference contrast microscopy, the extruded sperm were collected by overnight incubation at room temperature with Restore Western Blot Stripping Buffer (Pierce Biotechnology, Rockford, IL), treated with blocker and then reprobed with a different antibody.

**Histology of mouse testis**

**Fixation of testes.** Bouin’s fixative was introduced by vascular perfusion through the heart (Sprando, 1990). The mouse was given an intraperitoneal injection of heparin (Sigma-Aldrich) at 130 IU/kg body weight and then anesthetized with isoflurane 15 min later. The left ventricle was punctured with a blunted 22-gauge needle attached to a saline reservoir 3 ft above the animal. The right ventricle was cut immediately to allow a constant outflow of perfusate. After the perfusate had cleared of blood (~5 min), the fixative was switched on and allowed to perfuse through the mouse for 30 min. The testes were then removed and each one cut into two longitudinal sections and further exposed overnight to the fixative at 4°C. After several 30-min washes with PBS, the testes were washed with 50% ethanol to remove residual picric acid.

**Immunohistochemistry.** The fixed testes were embedded in paraffin for immunohistochemistry or LR White (Electron Microscopy Sciences, Fort Washington, PA) for histological studies. The procedure for antibody labeling was as described previously (San Agustin et al., 2000) with the following modifications. Antigens were retrieved by microwave heating (Panasonic model NN-5408A at high setting) for 10 min in 10 mM sodium citrate, pH 6. Before blocking with normal swine serum (Dako), the sections were treated with a biotin blocking system (Dako), which eliminated the faint background staining in the interstitial cells. For detection of IFT88, ropporin, and Odf2, sections were incubated overnight at 4°C with anti-IFT88 (1:200), antiropporin (1:250), or anti-Odf2 (1:250). Biotinylated Fab(α’b)2 fragment of affinity-isolated swine anti-rabbit immunoglobulins (Dako) was then applied (diluted to 0.71 μg/ml with TBS, 1% BSA [fraction V, fatty acid-free; Calbiochem/EMD Biosciences, San Diego, CA]), followed by alkaline phosphatase-conjugated streptavidin (diluted to 0.90 μg/ml; Dako). Exposure time was 20 min for both. The sections were washed with TBS between incubations. Labeling was visualized using the BCIP/NBT/INT substrate system (Dako), with a reaction time of 20–30 min. The sections were counterstained with hematoxylin (5 min), cleared with saturated lithium carbonate (1 min), and mounted with Glycergel.

**Periodic acid-Schiff staining.** The solutions used in PAS staining were prepared according to Tran et al. (2008). LR White sections (3 μm) were immersed in periodic acid for 1 h. After a brief rinse with water, they were transferred to the Schiff reagent for 1 h, rinsed with three changes (10 min each) of the sulfite solution, washed in running water (30 min), counterstained with hematoxylin (20 min), cleared with saturated lithium carbonate (2 min), air dried, and mounted with Shandon mount (Shandon, Pittsburgh, PA).

**Electron microscopy.** Testes were fixed by vascular perfusion as described, except that 5% glutaraldehyde in 50 mM sodium cacodylate, pH 7.4, was used as fixative. After perfusion, the testes were diced into 1-mm cubes, fixed in glutaraldehyde for another hour, washed with cacodylate buffer at 4°C, and then postfixed with 1% osmium tetroxide in 50 mM sodium cacodylate buffer, pH 7.4, at room temperature for 1 h. The tissue pieces were then washed three times in fresh cacodylate buffer and stained overnight at 4°C with fresh 1% uranyl acetate. After washing with water, the tissues were dehydrated through a graded series of ethanol and transferred to propylene oxide and finally into a 50:50 (vol/vol) mixture of Embed 812 (Electron Microscopy Sciences, Hatfield, PA) and propylene oxide for overnight infiltration. They were then transferred through three changes of fresh resin (1 h each) and finally polymerized at 60°C for 48 h.

**Isolation of spermatogenic cells.** All containers used were either siliconized or made of polypropylene. The two testes of an adult male (at least 8 wk old) were excised, transferred to 5 ml of KRB complete in a small Petri dish, and decapsulated. KRB complete is made up of Krebs-Ringer bicarbonate buffer (Sigma-Aldrich) plus 1.3 mM CaCl2, 1 mM glutamine, and 10 ml/l each of essential and nonessential amino acids (GIBCO/Invitrogen, Grand Island, NY), pH 6.9. The testes were then placed in 10 ml of 1 mg/ml collagenase...
A (Roche Applied Science, Indianapolis, IN) in KRB complete and incubated at 34°C for 15 min in a shaking water bath to uncoil the seminiferous tubules. The uncoiled tubules were allowed to settle, and the supernatant (which contained interstitial cells) was discarded. The tubules were washed three times (10 ml each) with KRB complete and resuspended in 5 ml of KRB complete. They were then diced (Vannas scissors; Ted Pella, Redding, CA) and aspirated from time to time to release the spermatogenic cells. The cell suspension was allowed to settle for 1 min, the supernatant collected, and the process repeated. The two supernatants (∼10 ml) were pooled and centrifuged (1000 rpm, 10 min, RC-5B centrifuge, SS-34 rotor) to collect the cells.

The soft pellet of cells obtained was resuspended in 1 ml of KRB complete and gently aspirated. The suspension was then layered on top of a Percoll step gradient (15, 22, 30, 45, and 90%, 1 ml each) and centrifuged at 2500 rpm for 30 min (RC-5B centrifuge, SS-34 rotor). The Percoll solutions were made by diluting Percoll in KRB basic (KRB complete minus CaCl₂, glutamine, and essential and nonessential amino acids). The spermatogenic cells usually collected between the 30 and 45% Percoll layers and were then resuspended in KRB basic to the desired density.

**Fixation of spermatogenic cells.** An equal volume of 2× fixing solution (4% paraformaldehyde in PBS) was added to the cell suspension and the cells fixed for −5 h on ice with constant, gentle shaking. An equal volume of 2% paraformaldehyde and 0.2% Triton X-100 in PBS was then added and the incubation continued for 30 min to permeabilize and further fix the cells. About 100 μl of fixed, permeabilized cells was then pipetted onto coverslips coated with 0.1% polyethyleneimine, allowed to settle for 10 min, and washed three times with 250 μl of PBS. The coverslips were stored in 400 μl of PBS at 4°C.

**Immunofluorescence microscopy.** Coverslips containing the fixed spermatogenic cells were rinsed with TBS and endogenous biotin blocked (Dako) following manufacturer’s instructions. This was followed by 1-h incubation in a blocking buffer (Tris-buffered saline [TBS], 5% BSA fatty-acid free, 10% nonimmune goat serum). Overnight double antibody labeling at 4°C was performed using 1) a polyclonal rabbit antibody to IFT88 and 2) a mouse monoclonal anti–α-tubulin antibody as marker for microtubules, including those of the manchette and axoneme. The latter was prepared by first incubating at room temperature the monoclonal anti–α-tubulin antibody (1:1000) with the F(ab′)₂ fragments of a goat anti-mouse immunoglobulin G (IgG) conjugated to Alexa 568 in TBS plus 1% BSA for 2 h, followed by 1-h incubation with normal mouse serum at 10% final concentration (to saturate the binding sites of the goat anti-mouse IgG F(ab′)₂ fragments). The rabbit anti-IFT88 antibody was added to this mixture just before use. On the next day the coverslips were rinsed three times in 250 μl of 1/5 blocking buffer. They were then incubated for 20 min with a 1:800 dilution of biotin-conjugated F(ab′)₂ fragment of goat anti-rabbit IgG in TBS and 1% BSA, rinsed three times with 250 μl of 1/5 blocking buffer, and then incubated for 20 min with a 1:400 dilution of streptavidin-conjugated Alexa Fluor 488 in TBS plus 0.5% BSA. After rinsing three times in TBS, the coverslips were counterstained with 0.5 μM TOTO-3 in TBS for 5 min, rinsed three times with TBS, and mounted using Prolong antifade. The fluorescent antibody and streptavidin conjugates, TOTO-3, and Prolong antifade were all purchased from Molecular Probes/Invitrogen (Eugene, OR).

Epididymal sperm were collected and fixed as described. After the 2-h fixation, they were permeabilized by mixing with an equal volume of fixative containing 0.1% SDS and left on ice for 15 min. About 200 μl of the permeabilized sperm was placed on a Super- frost Plus microscope slide (Fisher Scientific, Waltham, MA) and allowed to settle for 10 min. Slides were then washed with PBS, blocked with 1% BSA in TBS/Tween (TBST), and incubated with anti-acetylated α-tubulin antibody 6-11B-1 and anti-IFT88 antibody overnight at 4°C. Labeling by 6-11B-1 was visualized with Alexa Fluor 568 secondary antibody to mouse IgG; anti-IFT88 labeling was visualized with Alexa Fluor 488 secondary antibody to rabbit IgG (both secondary antibodies from Life Technologies, Carlsbad, CA). Between antibody incubations, slides were washed with 1% BSA in TBST. Nuclei were stained with 4′,6-diamidino-2-phenylindole dihydrochloride (DAPI; Invitrogen). Slides were mounted with coverslips using Prolong Gold Antifade (Life Technologies).

**Counting of germ-cell population in seminiferous tubule cross-sections.**

The procedure described by Oakberg (1956) was followed. Measurements were performed on 3-μm-thick LR White sections stained with PAS and hematoxylin. A crude value for the population of a particular germ cell in the tubule cross-section was obtained by counting all of their visible nuclei. The final, corrected count was obtained using the formula $F = C(t(1 + d))/t$, where $F$ is the final count, $C$ is the crude count, $t$ is the section thickness, and $d$ is the nuclear diameter. Nuclear diameters were measured using an ocular micrometer and then averaged.

**Detection of apoptotic cells.**

Testes were fixed with Bouin’s fixative as described and embedded in paraffin. Cut sections, typically 6 μm thick, were dewaxed in toluene and rehydrated with a graded series of isopropanol. The sections were transferred to 500 ml of 10 mM sodium citrate, pH 6.5, and heated for 5 min in a microwave oven (high setting). This partially disrupts the formaldehyde cross-links formed during the fixation process, thereby permeabilizing the sections. The permeabilized sections were cooled to room temperature with running tap water and then analyzed for apoptotic cells using the In Situ Cell Death Detection Kit, AP (Roche Molecular Biochemicals, Indianapolis, IN). The protocol recommended in the kit was followed, except that permeabilization was accomplished as described. The TdT enzyme stock solution was diluted 1:5 with the TUNEL dilution buffer (30 mM Tris-HCl, 140 mM sodium cacodylate, 1 mM CoCl₂, pH 6.6). The kit used fluorescein-labeled nucleotides as substrates for the TUNEL reaction. The sections were blocked (TBS containing 5% BSA, 20% normal rabbit serum) and the fluorescein-tagged, newly synthesized DNA products in apoptotic cells were then reacted with alkaline phosphatase-conjugated anti-fluorescein antibody. Antibody binding was visualized using a fuchsin substrate system (Dako). Color (magenta) was allowed to develop for 10 min, after which the sections were rinsed in water, counterstained with hematoxylin (5 min), cleared with saturated lithium carbonate solution (1 min), and mounted with Glycergel.

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