Mutations in a member of the Ras superfamily of small GTP-binding proteins causes Bardet-Biedl syndrome

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RAB, ADP-ribosylation factors (ARFs) and ARF-like (ARL) proteins belong to the Ras superfamily of small GTP-binding proteins and are essential for various membrane-associated intracellular trafficking processes^{1,2}. None of the ~50 known members of this family are linked to human disease. Using a bioinformatic screen for ciliary genes in combination with mutational analyses, we identified *ARL6* as the gene underlying Bardet-Biedl syndrome type 3, a multisystemic disorder characterized by obesity, blindness, polydactyly, renal abnormalities and cognitive impairment^{3,4}. We uncovered four different homozygous substitutions in *ARL6* in four unrelated families affected with Bardet-Biedl syndrome, two of which disrupt a threonine residue important for GTP binding⁵ and function⁵⁻⁷ of several related small GTP-binding proteins. Analysis of the *Caenorhabditis elegans* *ARL6* homolog indicates that it is specifically expressed in ciliated cells, and that, in addition to the postulated cytoplasmic functions of ARL proteins, it undergoes intraflagellar transport. These findings implicate a small GTP-binding protein in ciliary transport and the pathogenesis of a pleiotropic disorder.

Cilia and flagella are ancient, evolutionarily conserved eukaryotic organelles that project from cells and have been adapted by organisms to carry out diverse biological functions⁸. The assembly, maintenance and function of cilia and flagella depend on intraflagellar transport (IFT), and defects in this microtubule-based transport process and the function of cilia are associated with several human diseases, including Bardet-Biedl syndrome (BBS)⁸⁻¹⁰. Genes underlying seven of the eight loci known to be associated with BBS have been identified^{4,11}; only the gene mutated in BBS type 3 (called *BBS3*), previously mapped to 3p12 (refs. 12,13), remained unidentified. BBS is thought to result largely from ciliary dysfunction, because loss-of-function mutations in

C. elegans *bbs-7* and *bbs-8* compromise cilia structure and function¹⁴ and RNA interference of *Chlamydomonas* *BBS5* results in the loss of flagella¹¹. Notably, all known *C. elegans* *bbs* genes are expressed exclusively in cells with cilia, owing to the presence of a DAF-19 RFX transcription factor binding site (X box) in their promoters^{10,11}. We hypothesized that the *C. elegans* ortholog of human *BBS3* would also contain this regulatory element, which would allow us to identify candidates from the >90 genes that map to the *BBS3* critical interval^{12,13,15}. We generated a consensus X-box sequence from a training set of 14 *C. elegans* genes containing X boxes that are known to be strictly expressed in ciliated cells and used them to scan the *C. elegans* genome. We identified 368 genes with an X-box sequence within 1.5 kb of the start codon, 168 of which had a *bona fide* human ortholog (E value $\leq 10^{-6}$); three of these fell in the *BBS3* critical interval (Fig. 1a). The first gene, *ESRRBL1*, is probably the human ortholog of *C. elegans* *che-13*. *che-13* is expressed exclusively in ciliated neurons and has an important role in IFT¹⁶, and so *ESRRBL1* was an excellent candidate for *BBS3*. The second gene encodes the hypothetical protein DKFZp761H079, a member of the ARL family of small GTP-binding proteins^{1,2}. Its sequence is closely related to that of ARL2, and we called it ARL2-like protein 1 (ARL2L1). The third gene encodes ARL6, another ARL family member¹⁷.

To assess the likelihood that *ARL6* or *ARL2L1* is *BBS3*, we determined the expression patterns of their *C. elegans* orthologs (*arl-6* and predicted gene Y37E3.5, respectively). We generated transgenic lines expressing promoter-green fluorescent protein (GFP) fusion constructs (*arl-6p::gfp* or Y37E3.5p::gfp) and analyzed GFP fluorescence. Like the *che-13* promoter, the 5' untranslated regions (UTRs) of *arl-6* and Y37E3.5 directed expression to a small subset of sensory cells that are ciliated (Fig. 1b,c). For both transgenes, we observed GFP signals in the multiple ciliated amphid neurons in the head and both ciliated phasmid neurons (PHA and PHB) in the

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tail (Fig. 1b,c), as previously reported for other genes associated with BBS^{10,11}. We also detected expression in other ciliated sensory neurons, including the inner and outer labial neurons (Fig. 1b,c) and male tail ray neurons (data not shown). In the case of the *Y37E3.5p::gfp* transgene, we also detected GFP fluorescence in the midbody PDE ciliated neuron and PQR ciliated tail neuron.

Figure 1 Positions and expression analysis of candidate genes in the *BBS3* critical region. **(a)** The *BBS3* critical region on chromosome 3 is flanked by the markers *D3S1302* and *D3S1595* (refs. 12,13), although a recombination in one of the pedigrees suggests that the qter marker is actually *D3S1753* (ref. 12). The relative positions of the three candidate genes in the *BBS3* critical region are shown along with information relating to their *C. elegans* homologs. **(b,c)** Expression of *arl-6* and *Y37E3.5* in *C. elegans*. Transcriptional GFP reporters *arl-6p::gfp* **(b)** and *Y37E3.5p::gfp* **(c)** were expressed in the ciliated neurons of the worm head (e.g., amphid and inner and outer labial neurons) and worm tail (phasmids). Shown in the panels are differential interference contrast (DIC), GFP fluorescence and merged images of the whole worm, as well as higher magnification images of the head and tail regions for worms expressing *arl-6p::gfp*. The ciliated head and phasmid neurons of the tail are indicated by square brackets and arrowheads, respectively. The fluorescent signals along the midbody of worms expressing *arl-6p::gfp* is due to nonspecific autofluorescence of gut granules.

As the expression patterns of *C. elegans che-13*, *arl-6* and *Y37E3.5* are essentially identical to those of the five known *C. elegans* orthologs of genes associated with BBS^{10,11}, we examined their human orthologs for mutations in families with BBS. We sequenced the complete reading frames and exon-intron boundaries of *ESRRBL1* and *ARL2L1* in DNA of an affected individual from a well-characterized family from Newfoundland with BBS3 (NF-B2; ref. 13) but found no pathogenic mutations. However, we did identify a homozygous missense mutation 859G→C (resulting in the amino acid substitution G169A) in *ARL6* in this individual (Figs. 2 and 3a). The mutation segregated with the BBS phenotype in family NF-B2 (Fig. 2) and was absent from 100 control chromosomes. We next examined the genotypes of all available consanguineous families with respect to the eight *BBS* loci. The affected individual in a Saudi Arabian family (KK29) was homozygous with respect to markers across the *BBS3* critical interval, and association of the disease with all other loci was excluded in this family. Sequencing and subsequent segregation in this family showed that the affected individuals, but none of the unaffected relatives, were homozygous with respect to the mutation 445C→T, resulting in the nonconservative amino acid change T31M (Figs. 2 and 3a). To investigate further whether *ARL6* is *BBS3* and to determine its contribution to BBS, we expanded our mutational analyses to a multiethnic cohort of 230 families with BBS and identified two more families with mutations that are probably pathogenic. In North American family AR390, of suspected consanguinity, the affected individual, but none of the unaffected relatives, was homozygous with respect to the mutation 862T→G, resulting in the nonconservative amino acid change L170W (Figs. 2 and 3a). In addition, we identified an Irish consanguineous family, PB140, who carried the homozygous mutation 445C→G affecting the Thr31 residue (resulting in the amino acid substitution T31R; Figs. 2 and 3a). None of these changes were detected in 184 ethnically matched control chromosomes.

Mutations in *ARL6* segregate with BBS in four independent families, indicating that *ARL6* is *BBS3*. This idea is further supported by comparison of the amino acid sequences of *ARL6* from divergent organisms (Supplementary Fig. 1 online). Gly169 is invariant, and the

residues Thr31 and Leu170 are highly conserved: residue 31 is an amino acid with a hydroxyl side chain (threonine favored strongly over serine), and residue 170 is leucine (except for one valine) in 12 sequences (Supplementary Fig. 1 online). Of note, residue Thr31 lies in the highly conserved P loop of GTP-binding proteins (Fig. 3b). The analogous T31N mutation in ARF1 completely abrogates GTP binding *in vitro*⁵. Furthermore, overexpression of several small GTP-binding proteins mutated at the corresponding Thr31 residue produces cellular phenotypes consistent with loss of, or reduced, function. Therefore, such overexpression has been used extensively to study the functions of various ARF-ARL family members *in vivo*, including ARF1, ARF5, ARF6, ARL1, ARL3 and ARL7 (refs. 5–7,18–20). To determine the potential impact of the amino acid changes that we identified, we modeled ARL6 onto the structure of a homolog²¹, ARF6, which is 43% identical and 62% similar in amino acid sequence. All three altered residues cluster near, or are part of, the GTP binding site (Fig. 3c,d) and probably affect the GTP-binding activity and thus the function of ARL6. Taken together, these findings further indicate that the observed amino acid substitutions in ARL6 are pathogenic.

There is increasing evidence that the transmission of BBS is complex in some families, with a third mutation at a second locus acting as a modifier of either penetrance or expressivity^{4,22,23}. We identified a family, NF-B10, in which two affected sisters are homozygous with respect to the mutation 1179T→G in *BBS1* (resulting in the amino acid substitution M390R); one of them is also heterozygous with respect to the mutation 859G→C in *BBS3* (Supplementary Fig. 1 online). Clinical assessment indicated that the sister with three mutations is more severely affected with respect to some clinical features of BBS (Table 1), suggesting that the additional mutation in *BBS3* may act as a modifier.

RAB, ARF and ARL proteins are implicated in various aspects of biomembrane trafficking^{1,2}, although the function of ARL6 has not been explored in much detail. Hemagglutinin-tagged ARL6 is predominantly cytosolic, and associates with membranes *in vitro* in the presence of the nonhydrolyzable GTP analog GTPγS¹⁷. ARL6 interacts with the β subunit of the heterotrimeric protein import channel SEC61 (ref. 17), but the physiological relevance of this finding is unclear. To shed light on the function of ARL6, we generated *C. elegans* transgenic lines expressing GFP-tagged ARL-6. Analysis of these lines showed that ARL-6-GFP is cytosolic, as it was found in the cell bodies, dendrites, transition zone regions (akin to basal bodies found at the base of cilia) and ciliated structures in sensory neurons (Fig. 4a). Its cell localization is

Figure 3 The residues mutated in ARL6 cluster in or close to the GTP binding site in a structure modeled after ARF6. (a) Schematic of the gene *ARL6* showing the locations of mutations found in affected individuals from four families with BBS3. Exons are shown as shaded boxes. Mutations refer to transcript that includes exon 2a but not exon 2b. (b) Sequence conservation of the nucleotide binding site (P loop) in small GTP-binding proteins. Thr31 of HsARL6 is marked by an asterisk. (c,d) Homology modeling of ARL6 based on the crystal structure of ARF6 (ref. 21) shows the putative structure of ARL6. Amino acids that hydrogen-bond with the nucleotide GTPγS (shaded orange) are shown in blue. Amino acid changes resulting from the mutations identified in the families with BBS3 are shown in red.

therefore similar to that of GFP alone expressed in sensory neurons (Supplementary Video 1 online). To analyze whether ARL-6 may participate in dynamic, intracellular trafficking processes, we analyzed the transgenic lines by time-lapse microscopy. We observed that the GFP-tagged protein underwent IFT in the ciliary axoneme (Fig. 4b and Supplementary Videos 1–6 online). The movement of ARL-6-GFP occurred in both retrograde and anterograde directions along the cilium at 0.82 and 0.64 $\mu\text{m s}^{-1}$, respectively. These rates are comparable to those of other dynein- and kinesin-associated IFT proteins⁸.

Our finding that *BBS3* encodes a protein, ARL6, that is expressed specifically in ciliated cells and undergoes IFT in *C. elegans* has several important implications. ARL6 is the first BBS-associated protein

Table 1 Comparison of clinical features of two affected sisters in family NF-B10 with BBS

Clinical feature	PID 9	PID 22
Age at registration of blindness (y)	28	27
Maximal BMI (kg m^{-2})	45.8	34.5
Height (cm)	155	156
Chronic renal failure	Mild (onset at age 62 y)	None at age 41.8 y
Diabetes	Onset at age 31 y, on insulin since age 33 y	Not diabetic at age 41.8 y
Age at onset hypertension (y)	34	27
Polydactyly	None	None
Age at cholecystectomy (y)	36	None at age 41.8 y
Irritable bowel syndrome	Yes	No
Asthma	Yes, requiring a couple of hospital admissions	No
Psychiatric illness	Chronic depression, requiring medication	None
Renal stones	Age 51 y	None
Arnold Chiari malformation	Yes, requiring surgery	No
Idiopathic edema of upper and lower limbs	age 44 y	None
Dysfunctional uterine bleeding	Yes	No

Both individuals PID 9 and PID 22 are homozygous with respect to the mutation 1179T→G in *BBS1*; individual PID 9 is also heterozygous with respect to the mutation 859G→C in *BBS3*.

Figure 4 Localization and IFT of ARL-6 in the ciliated neurons of *C. elegans*. (a) Differential interference contrast (DIC), GFP fluorescence, and merged images of the head and tail regions of worms expressing a GFP-tagged ARL-6 fusion protein. ARL-6-GFP localized at the base of cilia in structures related to basal bodies (transition zones; Tz) and along the ciliary axonemes (Cil). Signals are also observed along the dendrites (Den) and axons (not shown) and within the main cell body (Cb). The cell bodies are shown only for the ciliated tail neurons. (b) Kymographs showing ARL-6-GFP particle movement. Scale bar is shown below each panel, and three time-frames (every 1,000 ms) show the movement of one particle (arrowheads) in the anterograde direction, away from the transition zones (intense staining on the left in each panel).

to be identified whose evolutionarily conserved sequence immediately suggests a cellular function, namely that of a small GTP-binding protein involved in membrane-associated intracellular trafficking processes¹. ARL6 is also the first known member of the RAB-SAR-ARF-ARL family of small GTP-binding proteins to be associated with an inherited human disease. The link between ARL6 and IFT extends the roles of this diverse protein family to include trafficking not only in the cytosol proper, but also in the ciliary axoneme. Given that BBS4 assists the targeting of a centrosomal protein, PCM1, in a dynein-dependent manner²⁴, and that other *C. elegans* BBS-associated proteins undergo IFT¹⁴, we propose that the BBS-associated proteins generally share a common function related to intracellular trafficking processes, such as the transport of cellular components (e.g., vesicle-associated or other proteins) to the centrosome, to the basal body and in the ciliary apparatus. Such a shared cellular function for BBS-associated proteins is expected, as the phenotypes associated with mutations at different BBS loci are indistinguishable⁴. Other small GTP-binding proteins, such as ARL2L1 identified in this study, may have similar cilia-specific functions. *Drosophila melanogaster* ARL3 and ARL6 are expressed specifically in chemo- and mechanosensory ciliated cells²⁵, and *Leishmania* ARL3 is required for flagellar integrity⁶. Similarly, mammalian ARL3 localizes to the photoreceptor connecting cilium in retinal rods and cones and, as such, could be implicated in retinitis pigmentosa²⁶. Several RAB GTPases have also been detected in the flagella of the green alga *Volvox carteri*²⁷. It will therefore be of particular interest to focus on the small GTP-binding protein family to determine how different members facilitate microtubule-based intracellular transport, and cilia- and flagella-related processes, especially in relation to human disorders that have a ciliary or neuronal component.

METHODS

Identification of BBS3 candidate genes. We created a profile Hidden Markov Model (HMM) using HMMER v1.8.4 (ref. 28) to identify *C. elegans* genes containing X-box regulatory sites in their upstream promoter regions. We constructed the profile HMM from a training set of 14 X box-containing genes whose expression is restricted to ciliated neurons: Y105E8A.5, F20D12.3, Y75B8A.12, T25F10.5, Y41G9A.1, T27B1.1, R31.3, F02D8.3, F38G1.1, F59C6.7, F33H1.1, K08D12.2, Y110A7A.20 and F40F9.1a. We used this profile HMM to scan the complete *C. elegans* genome (wormbase version WS110) and identified 368 genes with a consensus X-box sequence (raw score >14.0) present within 1.5 kb of the start codon. We downloaded from Ensembl a single FASTA format file containing the peptide sequences of all the *C. elegans* X box-containing genes that we identified using EnsMart. We downloaded a set of

unique *Homo sapiens* clusters from build 119 of UniGene from National Center for Biotechnology Information. Using Stand-Alone BLAST from National Center for Biotechnology Information, we compared the *C. elegans* protein sequences with the *H. sapiens* UniGene clusters by TBLASTN with an E-value cutoff set at 1×10^{-6} . For each *C. elegans* protein, we collected the top *H. sapiens* hit from the BLAST output and sorted these according to chromosome number and position. Three *H. sapiens* UniGene clusters fell within the BBS3 critical interval (Fig. 1a).

GFP fluorescence microscopy of *C. elegans* transcriptional and translational constructs. To produce transcriptional *gfp* constructs for the candidate *bbs-3* genes (*arl-6* and *Y37E3.5*), we used fusion PCR to introduce the 5' UTR sequence of each gene upstream of the full *gfp* coding sequence (including a nuclear localization signal sequence) and the *unc-54* 3' UTR¹⁰. The *arl-6p::gfp* construct contained 1,197 bp of the 5' UTR (and the first 14 bp of exon 1) and the *Y37E3.5p::gfp* construct contained 1,160 bp of the 5' UTR (and the first 14 bp of exon 1). We created translational *gfp* constructs for *bbs-3* (*arl-6p::gfp*) by fusing the entire exonic and intronic sequences of *arl-6* (including 1,197 bp of its 5' UTR) upstream of *gfp* (without a nuclear localization sequence) and the *unc-54* 3' UTR. These GFP transgenes were expressed as extrachromosomal arrays in *dpy-5(e907);Ex[dpy-5(+)]* worms, generated as described previously¹⁰. For live imaging, we immobilized worms (using 15 mM levamisole), mounted them on agarose pads and visualized them on a Zeiss Axioskop 2 + compound fluorescent microscope. Images and videos were captured using Northern Eclipse version 6.0 software.

Subjects. Individuals were diagnosed with BBS if they satisfied established criteria³. We assembled a large multiethnic cohort, comprising ~300 families with BBS. We obtained blood samples with consent, in accordance with protocols approved by the appropriate human subjects ethics committees at each participating institution, and extracted DNA using standard methods.

Mutational analysis of BBS3 candidate genes. We aligned the sequences of ARL6, *ESRRBL1* and ARL2L1 with the corresponding human genome sequences and determined the exon-intron boundaries. We identified sequences flanking all coding exons and used them to design primers (sequences available on request) that amplify the exons and exon-intron boundaries of each gene. PCR amplification products were purified, sequenced and analyzed as described²³.

Protein sequence analysis and homology modeling. We obtained ARL6 protein sequences from GenBank and aligned them using the default settings with ClustalX. *H. sapiens* ARF6 was manually aligned to ARL6 based on a previous report². Homology modeling of HsARL6 was done using Swiss-Model with HsARF6 as a template. The resulting ARL6 three-dimensional model was manipulated and rendered in PyMOL.

URLs. Online Mendelian Inheritance in Man is available at <http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=OMIM>. Wormbase is available

at <http://www.wormbase.org/>. Ensembl is available at <http://www.ensembl.org/>. Swiss-Model is available at <http://www.expasy.org/swissmod/SWISS-MODEL.html>. Protein Data Bank is available at <http://www.rcsb.org/pdb/>. The National Center for Biotechnology Information is available at <http://www.ncbi.nlm.nih.gov/>. Hidden Markov Model is available at <http://hmmer.wustl.edu/>. PyMOL is available at <http://www.pymol.org>.

Accession numbers. GenBank: *ESRRBL1*, NM_018010; *ARL2L1* (encoding hypothetical protein DKFZp761H079), NM_182896; *ARL6*, NM_032146. Protein Data Bank: HsARF6, 1HFV.

Note: Supplementary information is available on the Nature Genetics website.

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COMPETING INTERESTS STATEMENT

The authors declare that they have no competing financial interests.

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