The zebrafish mutant lbk/vam6 resembles human multi-systemic disorders caused by aberrant trafficking of endosomal vesicles

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The trafficking of intracellular vesicles is essential for a wide range of cellular processes and defects in this process have been implicated in a wide range of human diseases. We identify the zebrafish mutant lbk as a novel model for such disorders. lbk displays hypopigmentation of skin melanocytes and the retinal pigment epithelium (RPE), an absence of iridophore reflections, defects in internal organs (liver, intestine) as well as functional defects in vision and the innate immune system (macrophages). Positional cloning, an allele screen, rescue experiments and morpholino knock-down reveal a mutation in the zebrafish orthologue of the vam6/vps39 gene. Vam6p is part of the HOPS complex, which is essential for vesicle tethering and fusion. Affected cells in the lbk RPE, liver, intestine and macrophages display increased numbers and enlarged intracellular vesicles. Physiological and behavioural analyses reveal severe defects in visual ability in lbk mutants. The present study provides the first phenotypic description of a lack of vam6 gene function in a multicellular organism. lbk shares many of the characteristics of human diseases and suggests a novel disease gene for pathologies associated with defective vesicle transport, including the arthrogryposis-renal dysfunction-cholestasis (ARC) syndrome, the Hermansky-Pudlak syndrome, the Chediak-Higashi syndrome and the Griscelli syndrome.

KEY WORDS: Zebrafish development, Eye, Vision, Pigmentation, Liver, Vesicle trafficking, Lysosomes and lysosome-related organelles, Vam6p/Vps39

INTRODUCTION

Vesicle trafficking is essential for a wide range of cellular processes and defects in vesicle trafficking have been shown to underlie various human diseases. In the arthrogryposis-renal dysfunction-cholelasis (ARC) syndrome, for example, mutations in the VPS33B gene lead to a dysregulation of vesicle fusion, including lysosomes and late stage endosomes (Gissen et al., 2004) (OMIM #208085). Similarly, the Chediak-Higashi syndrome (CHS), Hermansky-Pudlak syndrome (HPS) and Griscelli syndrome (GS) have been linked to a number of proteins involved in the formation, trafficking and fusion of lysosome and lysosome-related organelles (OMIM #214500 (CHS), #213300 (HPS), #214450 (GS1), #607624 (GS2), #609227 (GS3)). These inherited human diseases are associated with hypopigmentation, prolonged bleeding and immunological defects. It has also been reported that affected individuals display visual defects, including photophobia, strabismus and nystagmus. Although mutations in several genes have been linked to these diseases, it has not yet been possible to identify the molecular defects in all cases.

Lysosome-related organelles are cell type-specific organelles that share several physiological properties with lysosomes (Raposo and Marks, 2002). They include organelles, such as melanosomes in skin and RPE melanocytes, lamellar bodies in type II lung epithelial cells, dense granules in platelets and MHC class II compartments in antigen-presenting cells, and cytotoxic granules of lymphocytes. Tethering and docking of lysosomes and lysosome-related organelles depends on the HOPS complex (homotypic fusion and vacuole protein sorting complex). This complex interacts with both SNAREs and Ypt7p/Rab7 (Collins et al., 2005; Price et al., 2000a; Price et al., 2000b; Sato et al., 2000; Wurmser et al., 2000) and is required for SNARE complex assembly (Stroupe et al., 2006). Furthermore, the HOPS complex seems to be important for early-late endosome transition in the Rab-conversion model in mammalian cells (Rink et al., 2005). The HOPS complex comprises the class C Vps protein Vam5p/Vps33p, the protein defective in ARC syndrome; the class C proteins Vam1p/Vps11p, Vam9p/Vps16p and Vam8p/Vps18p; and two class B Vps proteins, Vam6p/Vps39p and Vam2p/Vps41p (Rieder and Emr, 1997; Seals et al., 2000; Wurmser et al., 2000) (reviewed by Bowers and Stevens, 2005).

Vam6p/Vps39p was shown to be required for vacuolar protein sorting in yeast. Yeast Vps39p-null mutants exhibit highly fragmented vacuolar morphogenesis and mutant cells accumulate numerous vesicular structures scattered throughout the cytoplasm (Nakamura et al., 1997). Similarly, overexpression studies in cultured HEK cells have implicated Vam6p in the clustering and fusion of lysosomes and late endosomes (Caplan et al., 2001) and blocking Vam6p function with antibodies suggests a possible role in sperm cells during their interactions with the zona pellucida in mice (Brahmaraju et al., 2004).

We have identified a zebrafish mutant (leberknödel; lbk) that carries a mutation in the zebrafish homologue of Vam6p/Vps39p and displays phenotypes similar to those observed in humans suffering from ARC syndrome, CHS, HPS and GS. Our studies reveal multi-systemic defects in lbk, including a hypopigmentation of skin melanocytes and the retinal pigment epithelium (RPE). Moreover, lbk displays defects in internal organs (liver, intestine) and the innate
immune system. Affected cells display increased numbers and enlarged intracellular vesicles. Physiological and behavioural analyses of visual function in \( lbk \) uncovered a reduced visual ability. These analyses suggest that \( vam6p/Vps39p \) has essential functions in a range of tissues during zebrafish development. Notably, no animal model to study the function of \( vam6p/Vps39p \) in vivo has been reported to date.

### MATERIALS AND METHODS

#### Fish maintenance and strains

Zebrafish (\textit{Danio rerio}) were kept under standard conditions. Strains used in this study were: Tü (Haffter et al., 1996), WIK (Dahm et al., 2005) and \( lbk+/- \) (Rinner et al., 2005b). Staging of embryos in hours (hpf) and days post-fertilization (dpf) was carried out as previously described (Kimmel et al., 1995).

#### Transplantations

Transplantations were performed as described (Ho and Kane, 1990). Mosaic animals were generated by transplanting 30-40 cells from 3-4 hpf wild-type Tü embryos into the animal pole region, including prospective ectodermal domains, such as the eye and neural crest domains (Kimmel et al., 1995), of age-matched embryos obtained by crossing two \( lbk+/- \) carriers. Larvae displaying a clear \( lbk+/- \) phenotype were scored for the presence of skin melanocytes and RPE cells displaying wild-type levels of melanin.

#### Histology, immunohistochemistry and TUNEL assay

For histological analyses by light microscopy (LM), larvae were fixed in 4% paraformaldehyde (PFA) in PBS (pH 7.2) at 4°C overnight and washed three times in PBS. The embryos were dehydrated in a standard ethanol series, infiltrated and embedded in Technovit 7100 (Heraeus Kulzer, Germany) for at least 2 hours, embedded in OCT TissueTek (Jung-Leica; Tissue Freezing Medium) and frozen in liquid nitrogen (N2). Sections (20 m) were collected on coated slot grids, stained with uranyl acetate and lead citrate and viewed in a Philips CM 10 electron microscope.

For immunohistochemistry, larvae were fixed in 4% PFA in PBS (pH 7.2) for 45 minutes at room temperature, cryoprotected in 30% sucrose for at least 2 hours, embedded in OCT TissueTek (Jung-Leica; Tissue Freezing Medium) and frozen in liquid nitrogen (N2). Sections (20 m) were cut with a glass knife and mounted on Superfrost Plus slides (Microm International, Switzerland). The sections were subsequently washed with 1% aqueous uranyl acetate for 1 hour at 4°C, dehydrated through a graded series of ethanol, infiltrated with ethanol/resin mixtures and embedded in Epon (using glycidether 100 from Roth, Karlsruhe). Ultra-thin sections were collected on coated slot grids, stained with uranyl acetate and lead citrate and viewed in a Philips CM 10 electron microscope.

For confocal laser scanning microscopy (CLSM), larvae fixed in 4% PFA in PBS (pH 7.2) were blocked with a solution of 20% normal goat serum and 2% bovine serum albumin in PBS containing 0.3% Triton X-100 (PBST) for 1 hour. Sections were then incubated for 2 hours at 4°C in the primary antibody. Primary antibodies used in this study were anti-glutamine synthetase (1:700; Chemicon, Temecula, USA) and anti-rodopsine (1:200; MilanAnalytica, LaRoche, Switzerland). Antibodies were diluted in PBST. After three washes in PBST, sections were incubated in Alexa546-coupled anti-mouse secondary antibody (1:500; Molecular Probes/Invitrogen, Eugene, OR) for 1 hour, washed three times in PBST, mounted in glycerol and analysed with an Axioscope fluorescence microscope as described above. Apoptosis was detected using the TUNEL (TdT-mediated dUTP nick-end labelling) method on 35 m cryosections. TUNEL staining was performed using the Cell Death Kit (Roche Molecular Biochemicals) according to the manufacturer’s protocol. Oil red O in dextrin was used to visualise lipid-containing vesicles in cryosections (20 m). Confocal laser scanning microscopy (CLSM) was performed with an LSM510 (Zeiss) and 25× PlanNeofluar and 63× PlanAPOCHROMAT oil immersion objectives (Zeiss).

#### Genetic mapping of \( lbk \) and radiation hybrid mapping of \( vam6 \)

Map crosses were set up between heterogeneous \( lbk+/- \) (Tü background) and wild-type WIK zebrafish. The offspring from these crosses were inbred and homozygous \( lbk+/- \) and sibling F2 progeny were collected and their DNA extracted. Bulked segregant analysis on 48 \( lbk+/- \) larvae and 48 siblings, respectively, was carried out using 192 simple sequence length polymorphisms (SSLP markers) distributed over the entire genome (Geisler, 2002). Further fine mapping was performed using the total DNA of single homozygous mutant larvae. DNA extraction andPCR were performed as described (Geisler, 2002). The zebrafish \( vam6 \) gene was radiation hybrid (RH) mapped on the Goodfellow T51 RH panel as described (Geisler, 2002) using six independent primer pairs (Table 1). PCRs for RH mapping were carried out independently in triplicate.

#### Cloning of zebrafish \( vam6 \)

A putative zebrafish \( vam6 \) gene was identified via a homology search of the Zv5 Ensembl database (http://www.ensembl.org/Danio_rerio) using the human Vam6p protein sequence as query. Total RNA was isolated from adult zebrafish eyes using the RNAeasy kit (QIAGEN, Switzerland) according to the manufacturer’s instructions. cDNA was synthesised using the SuperScript III Reverse Transcriptase Kit (Invitrogen, Switzerland) as described in the manual. The zebrafish \( vam6 \) cdns was cloned using primers designed against the start and stop regions of the \( vam6 \) cDNA (Table 2), respectively, using homology with the human VAM6 gene (VPS39 — Human Gene Nomenclature Database). To obtain 5’ sequences of the \( vam6 \) cDNA, RACE-PCR was carried out with the SMART RACE cDNA Amplification Kit (Clontech, Switzerland). For RACE primers used to amplify 5’ regions of the \( vam6 \) cDNA, see Table 2. PCR products were cloned into the pCRII-Vector (Clonetech), sequenced with T7 and SP6 primers and submitted to GenBank (Accession Number, EF446162).

#### Allele screen

Heterozygous \( lbk+/- \) carriers were crossed to a collection of ENU-mutated Tü zebrafish and the offspring screened for phenotypes equivalent to that of \( lbk+/- \), indicating non-complementation of the novel allele with the \( lbk \) mutation. Total genomic DNA from 5 dpf larvae showing an \( lbk \)-like phenotype was extracted, all 24 exons of the \( vam6 \) gene were amplified and the PCR products sequenced (Table 3). All PCR and sequencing reactions were performed independently in triplicate.

#### Optokinetic response measurements and electroretinogram recordings

Optokinetic response (OKR) measurements were performed as described previously (Rinner et al., 2005b). Briefly, single larvae were placed in the centre of a Petri dish (35 mm diameter) filled with 3% pre-warmed (28°C) methylcellulose facing with their right eye a screen at an apparent distance of

<table>
<thead>
<tr>
<th>RH primer pair</th>
<th>Sense primer</th>
<th>Antisense primer</th>
</tr>
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<tbody>
<tr>
<td>1</td>
<td>ATTTGTGTCGGACAGGTATAC</td>
<td>TAGGATTTAGTGTTTGACGCTC</td>
</tr>
<tr>
<td>2</td>
<td>TGTCAGCTGCTGCCAGCTGATCT</td>
<td>TAGGAGTTAGGTGGGGAAGGCTC</td>
</tr>
<tr>
<td>3</td>
<td>AGACGTTCAGATCTGCTGAGGG</td>
<td>TGCCTCGACATTCTCTAGTGC</td>
</tr>
<tr>
<td>4</td>
<td>GAGGAGCTTGTGCCTGCTGCTGTC</td>
<td>TCTCGGCGTCTTGTAGTGG</td>
</tr>
<tr>
<td>5</td>
<td>AACTCCGCTGTTGAGGCGG</td>
<td>TCTCGGCGTCTTGTAGTGG</td>
</tr>
<tr>
<td>6</td>
<td>TGCTCTCCCTCTCGTACTCC</td>
<td>TCTCGGCGTCTTGTAGTGG</td>
</tr>
</tbody>
</table>
4.65 cm within their visual field. Moving sine-wave gratings were projected by a HP vp6111 projector onto the screen. The projection size on the screen was 8 x 6 cm, subtending a visual angle of 65.6° horizontally and 53.1° vertically. Eye movements were recorded by an infrared-sensitive CCD camera triggered by the visual stimulation. A custom developed programme based on the LabView IMAQ software (National Instruments, version 5.1) was used to control the stimulation and the camera as well as to analyse the resulting images. Contrast sensitivity functions for mutant and wild-type larvae were measured by the eye velocity as a function of the contrast of the moving grating. Statistics were derived as described previously (Rinner et al., 2005a) by bootstrap sampling of differences between randomly drawn samples (n=10,000) from a pooled data set of wild-type and mutant larvae. The significance level chosen was at 5% for all experiments.

For recordings of electroretinograms (ERGs) (Makhankov et al., 2004), 5 dpf larvae were dark-adapted for at least 30 minutes prior to positioning them in the recording chamber. The recording chamber was shielded from interference from external sources of electromagnetic radiation by placing it inside a tight Faraday cage. Each larva was placed on its side on the surface of a moist sponge with E3 medium (5 mM NaCl, 0.17 mM KCl, 0.33 mM CaCl2; and 0.33 mM MgSO4) and paralyzed by directly adding a drop of the muscle relaxant Esmeron (0.8 mg/ml; Organon Teknika, Eppelheim, Germany) onto the larva. An Ag/AgCl electrode system was used to record the ERG responses. The recording electrode was positioned on the centre of the cornea. The reference Ag/AgCl pellet was placed under the body of the larva. All pre-recording steps were carried out under dim red light illumination. An additional 5-minute period in complete darkness was chosen to adapt the larva to dark prior to measurements. White light stimuli (100 ms) were used to elicit ERG responses with interstimulus intervals of 5 seconds. Light stimuli were fixated at five relatively different light intensities ranging from 2 lux (OD 5) to 20,000 lux (OD 0). Unattenuated light stimulus intensities were measured over the head of the larvae using a light meter (Tektronix J17, Texas Instruments, USA) and found to account for 3100 lux (optical density (OD) equal to 0 log units). Different light intensities were adjusted using neutral density filter wheels. A virtual instrument (VI) under NI LabVIEW 5.1 was developed for use in all experiments. Sampling was carried out in buffered acquisition mode with a sampling rate of 1000 Hz. The resulting ERGs, as the corneal sum field potentials of the retina in response to light, are shaped like in other vertebrates: a small negative deflection, termed the a-wave, which reflects photoreceptor activation, is followed by a stronger positive deflection, the b-wave, which reflects second order neuron activation. The small a-wave is often masked by the much larger b-wave because of interference. Therefore, in contrast to the b-wave, the a-wave is not a robust measure in larval zebrafish. As a consequence, the b-wave was taken as an indirect measure of outer retina activation. For rescue experiments, a light intensity of 6000 lux was chosen to elicit ERG responses in sibling and lbk−/− larvae. ERG responses obtained from one larva were averaged three to seven times depending on signal-to-noise ratio. Statistical analysis was performed using GraphPad Prism4 (GraphPad Software, San Diego, CA) software and graphs were generated using Origin v.7 (OriginLab, Northampton, MA).

**Bacteria injections**
Salmonella enterica serovar Typhimurium expressing DsRed (van der Sar et al., 2003) were injected with a FemtoJet injection apparatus (Eppendorf) at a constant pressure of 80 hPa for 0.5 seconds into the common cardinal vein of 48 hpf larvae embedded in low-melting point agarose as previously described (van der Sar et al., 2003). To ensure delivery of equal amounts of bacteria, mutant and sibling larvae were injected with the same injection capillary and the injection of the bacteria was monitored under a fluorescence stereo-microscope (Zeiss). For bacterial counts, larvae were macerated in PBS and plated onto ampicillin-containing bacterial plates as described (van der Sar et al., 2003; van der Sar et al., 2006). Bacterial growth was assessed after 24 hours of incubation at 37°C with a fluorescence stereo-microscope.

**Rescue experiments**
The rescue was performed by injecting a pSGH2 vector containing the van6 cds under the control of a heat-shock promoter (Bajoghli et al., 2004) into one-cell stage embryos derived from a cross of two lbk−/− individuals. At 3 dpf, injected larvae were heat shocked at 38°C for 2 hours and subsequently transferred back to 28°C. The extent of the rescue was determined at 5 dpf by assessing the level of pigment in skin melanocytes and RPE and in ERG measurements.

**Morpholino knock-down**
A 25 nucleotide morpholino antisense oligonucleotide was designed against the ATG region of the van6 mRNA: GAACTGTTGTCGATCAGCGTGCAATCAAGG (downstream of STOP). The morpholino was injected using a pressure of 30 hPa for 4 msedents into one-cell stage embryos by direct injection of the bacteria was monitored under a fluorescence stereo-microscope. The significance level chosen was at 5% for all experiments.

**Detection of van6 expression**
For PCR analysis, total RNA was isolated from 100 wild-type zygotes and 4 hpf embryos, respectively, using the RNAeasy kit and cDNA synthesised using the SuperScriptIII Reverse Transcriptase kit. PCR was performed at an annealing temperature of 60°C for 35 cycles using primers listed in Table 2.

**Whole-mount in situ hybridisation**
Sence and antisense probes for in situ hybridisation were in vitro transcribed from a linearized pCRII-Vector containing the full-length van6 cds using SP6 and T7 polymerases, respectively, in the presence of digoxigenin-coupled nucleotides (DigRNA labelling kit; Roche Molecular Biochemicals, Switzerland). Transcripts were hydrolysed to obtain fragments of ~300-500 nucleotides in length. To prevent melanisation of skin melanocytes and the

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**Table 2. Primer pairs used for the cloning of the zebrafish van6 cDNA and detection of zebrafish van6 expression in zygotes and 24 hpf embryos**

<table>
<thead>
<tr>
<th>Experiment</th>
<th>Sense primer</th>
<th>Antisense primer</th>
</tr>
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<tbody>
<tr>
<td>Cloning of van6 cds</td>
<td>CAGGAGATCTACGGGTTTGG (upstream of START)</td>
<td>AAAGTCAGATCCAGACAGAAGAC (downstream of STOP)</td>
</tr>
<tr>
<td>Cloning of van6 cds</td>
<td>TGGTTTTCTGTCTGGTTTGG (upstream of START)</td>
<td>AAAGTCAGATCCAGACAGAAGAC (downstream of STOP)</td>
</tr>
<tr>
<td>5’ RACE PCR</td>
<td>CAGGATATAGGATTCTACACACAG</td>
<td>Universal primers as supplied with the SMART RACE cDNA Amplification Kit (Clontech, Switzerland)</td>
</tr>
<tr>
<td>Nested 5’ RACE PCR</td>
<td>GAGCGATGACATCTCCCGTAATTCG</td>
<td>Universal primers as supplied with the SMART RACE cDNA Amplification Kit (Clontech, Switzerland)</td>
</tr>
<tr>
<td>Detection of short van6 fragment by PCR</td>
<td>ATGATCGACGCGTACGAACAGGTC</td>
<td>GGGATGTCAGTTGAGG</td>
</tr>
<tr>
<td>Detection of long van6 fragment by PCR</td>
<td>CAGGAGATCTACGGGTTTGG</td>
<td>AAAGTCAGATCCAGACAGAAGAC</td>
</tr>
<tr>
<td>Detection of β-actin fragment by PCR</td>
<td>GAGCAAGTCTGTCATGTTGG</td>
<td>TTGCAGATGTGATGACC</td>
</tr>
<tr>
<td>Detection of pmel17 fragment by PCR</td>
<td>TTCATGGAATCACCAGATGTACC</td>
<td>GAACGTAGTGGATATACATAACCAAAGG</td>
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</table>
Table 3. Primer pairs used to amplify and sequence the 24 exons of the vam6 gene from total genomic DNA of 5 dpf larvae showing an lbk-like phenotype (lbk*) in the allele screen

<table>
<thead>
<tr>
<th>Exon</th>
<th>Sense primer</th>
<th>Antisense primer</th>
</tr>
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<tbody>
<tr>
<td>Exon 1</td>
<td>TGGGTGGTTCCTCGTGTTG</td>
<td>CAAATGAAAGTGAGAAAGAGCATGAC</td>
</tr>
<tr>
<td>Exons 2 to 5</td>
<td>TGACCCTGACATATTAGTACAC</td>
<td>TGTTGCAAGTGAAGATAAGTGC</td>
</tr>
<tr>
<td>Exons 6 and 7</td>
<td>ACTTTTCTCAGCATTGTCGTTCC</td>
<td>CAACGCTGATACCAACTAAGG</td>
</tr>
<tr>
<td>Exon 8</td>
<td>CGTCGTACACATTTTCTGTGAACG</td>
<td>CTGAAACCGGGGGGAAAAAG</td>
</tr>
<tr>
<td>Exon 9</td>
<td>TGCACACCTGTTGATCCAGCC</td>
<td>TTCAACGATCCTGGTGC</td>
</tr>
<tr>
<td>Exons 10 and 11</td>
<td>AAAAAGTGGTGCAGGTTTACGG</td>
<td>CAAACCGGTATCCTGGTG</td>
</tr>
<tr>
<td>Exons 12 and 13</td>
<td>TCTGGGTGTTTTCGTGTTGGCC</td>
<td>TTGGACAAAAGAGACAGAGG</td>
</tr>
<tr>
<td>Exon 14</td>
<td>TGCAACATGGTGGAAAGG</td>
<td>AAAACGGTTCAGCGCAAGAG</td>
</tr>
<tr>
<td>Exon 15</td>
<td>TGACTCCAGACCTGGAAACG</td>
<td>AAGAAATTGCAGGAAATAGG</td>
</tr>
<tr>
<td>Exons 16 and 17</td>
<td>CGAAGTCTACATGTCGTGAACC</td>
<td>ATCACTCCAGCTGTTG</td>
</tr>
<tr>
<td>Exons 18 and 19</td>
<td>GGAGACGTGCTCGGACGATG</td>
<td>ATCACTCCAGCTGTTG</td>
</tr>
<tr>
<td>Exon 20</td>
<td>CAACGCTGACATATTAGTACAC</td>
<td>GGATTTGTCATGACAGAAGG</td>
</tr>
<tr>
<td>Exon 21</td>
<td>GGTGAGCATTTTCTGTTGAAAGG</td>
<td>GGTCTCCTCATCCGAGAAG</td>
</tr>
<tr>
<td>Exon 22</td>
<td>TTGGACACATTCTTCTGTTGAAAGG</td>
<td>CAAATTGCCAGAAAAACCAAGGA</td>
</tr>
<tr>
<td>Exon 23</td>
<td>AAGAATGGCAGGAGCATATTAGCAG</td>
<td>TTTTTACCAGGATCAGG</td>
</tr>
<tr>
<td>Exon 24</td>
<td>GGATTCGTTGAGATGAGGCAGG</td>
<td>CATCCTAAGTGAAGGAGG</td>
</tr>
</tbody>
</table>

In cases where a primer pair was used to sequence more than one exon, intervening introns were also amplified and sequenced.

RESULTS

lbk mutants show hypopigmentation and hepatomegaly

The zebrafish mutant leberknödel (lbk) (Glass and Dahm, 2004), was identified in a largescale ENU mutagenesis screen (Glass and Dahm, 2004). External analyses revealed that, starting at 3 dpf, lbk–/– larvae show a pronounced hypopigmentation of skin melanocytes and the RPE. Instead of the evenly black melanocytes characteristic of wild-type larvae, lbk melanocytes show only scarce and patchy pigmentation (Fig. 1, see Fig. S1 in the supplementary material). It is interesting that in addition to melanosome-containing cells the iridophores are affected in lbk. Iridophores derive their reflective properties from vesicular organelles containing guanine crystals (Morrison and Frost-Mason, 1991). These light reflections are absent in lbk, both in the eye and body, at all stages of development (Fig. 1E-H), indicating that the biogenesis of these organelles is disrupted. We cannot, however, exclude the possibility that iridophores are missing altogether, although, in view of the overall lbk phenotype, this possibility appears unlikely. Further to the pigment cell phenotypes, starting at 4 dpf, the intestinal tract of lbk develops a brownish discoloration (Fig. 1A-F,J) and the liver becomes enlarged (Fig. 1I-J). lbk–/– larvae fail to develop swim bladders and die between 7 and 8 dpf.

Mapping and candidate gene approach identify a mutation in the vam6 gene as underlying the lbk phenotype

To identify the mutation underlying the lbk phenotype, we mapped the mutation on the zebrafish genome. Pooled DNA of 48 lbk–/– and 48 sibling 5 dpf larvae, respectively, was tested with 192 SSLP markers (Knapik et al., 1998) resulting in a linkage of the mutation to chromosome 17 between markers z22083 and z4053 (Fig. 1K). To narrow down the critical interval of the mutated locus, we tested additional SSLP markers located in this genomic region. Analysis of 282 meioses identified z9692 and z4053 as the closest SSLP markers (Fig. 1K).

As the phenotype observed in lbk suggests a defect in endosomal vesicle trafficking (see below), we performed RH mapping of several genes involved in this process. This approach resulted in a linkage of the zebrafish vam6 gene to the EST fc27c07.x1 on chromosome 17. This EST is located in close proximity to z9692 on the T51 RH map (Geisler et al., 1999). Given this close linkage, we cloned and sequenced the zebrafish vam6-coding sequence (cds), which has a length of 2628 bp corresponding to a protein of 875 amino acids (see Fig. S2 in the supplementary material).

Sequencing of pooled vam6 cDNA from lbk–/– larvae and phenotypically wild-type siblings, respectively, revealed a nonsense mutation at position 1066 of the vam6 cds (Fig. 1L), which segregated with the mutant phenotype. This point mutation was confirmed by sequencing exon 11 from genomic DNA of the larvae identified as recombinant in the fine mapping approach. All 15 individuals displaying the lbk phenotype were heterozygous or homozygous for the point mutation, whereas all phenotypically wild-type individuals were either heterozygous or homozygous for the wild-type allele (not shown). The putative truncated protein product lacks the C-terminal 520 amino acids, including the central clathrin homology (CLH) repeat domain, which has been demonstrated to play an essential role in the clustering and fusion of lysosomes (Caplan et al., 2001), and Ypt7p-interacting sequences (Wurmser et al., 2000) (Fig. 1M,N).

To identify additional lbk alleles and thus confirm that a mutation in vam6 underlies the lbk phenotype, we performed complementation analyses with described mutants displaying similar external phenotypes (Glass and Dahm, 2004; Kelsh et al., 1996). These analyses identified no additional lbk alleles. We therefore performed an allele screen by crossing heterozygous Tü zebrafish. Over 1000 genomes were screened for external phenotypes equivalent to that observed in lbk, indicating non-complementation of the novel allele with the lbk mutation. This approach identified a compound heterozygous larva (lbk+; see Fig. S3 in the supplementary material) harbouring in one allele of the vam6 gene the original C➔T exchange found in lbk and in the second allele a T➔A exchange at position 374 of the vam6 cds (Fig. 1O). The latter mutation results in the exchange of a
The lbk mutation acts cell autonomously

To investigate if the defect caused by the mutation in lbk is cell-autonomous, we performed blastomere transplantsations. Wild-type cells from 1000-cell stage embryos were transplanted into age-matched embryos obtained from a cross of two heterozygous lbk carriers. At 5 dpf, the resulting larvae were screened and larvae with a predominantly lbk−− mutant skin phenotype (as judged by the presence of strongly hypopigmented melanocytes) were investigated for the presence of melanocytes of wild-type appearance. As the lbk−− phenotype is fully penetrant such that mutants never display melanocytes with normal levels of pigmentation, melanocytes of wild-type appearance had to be derived from transplanted cells. In four independent experiments, a total of 26 lbk−− background larvae displaying melanocytes with wild-type pigmentation in their skin were identified (see Fig. S4 in...
the supplementary material). The development of transplanted wild-type cells into normally pigmented melanocytes in an lbk<sup>−/−</sup> background indicates that the lbk mutation acts cell autonomously.

**lbk larvae have a compromised RPE and are severely visually impaired**

Hypopigmented retinas often have additional defects and affected zebrafish were shown to display decreased visual ability (Schonthaler et al., 2005). Similarly, in humans several pathological conditions with RPE hypopigmentation are associated with more generalised ocular symptoms. This prompted us to examine the eye phenotype in lbk more closely. Histological analyses revealed that the overall patterning of the neural retina is normal (Fig. 2A-F). However, sections taken at different developmental stages revealed not only a severe hypopigmentation of the RPE, but also a progressive thickening (Fig. 2A-H) up to three times the thickness of the wild-type RPE (Fig. 2R). This thickening results from an increase of RPE cell size, rather than increased numbers of RPE cells (Fig. 2S). TEM analyses of lbk revealed substantially fewer and mostly immature and aberrantly shaped melanosomes (Fig. 2I-N,Q). As melanin is, however, present in the lbk<sup>−/−</sup> RPE and skin melanocytes (see Fig. S5 in the supplementary material), the mutation affects the biogenesis of melanosomes rather than melanin biosynthesis. This is supported by the observation that the number of mature melanosomes is substantially reduced in lbk (Fig. 2V).

In addition to optically shielding photoreceptor cells (PRCs), the RPE is essential for PRC function by phagocytosing shed PRC outer segments. Outer segments harbour the photopigments that allow the detection of light and the shedding of parts of these segments is an essential process in the maintenance of normal PRC function. The RPE cytoplasm in lbk is filled with numerous vesicles whose number increases between 5 and 7 dpf (Fig. 2I-L). Some of these vesicles contain thick membrane stacks reminiscent of the membrane discs in PRC outer segments (Fig. 2M,N), suggesting a defect in the fusion of endocytic vesicles with lysosomes.

To test this hypothesis, we used rhodopsin as a molecular marker for rod outer segments (Fig. 2O,P). lbk larvae (5 dpf) show an increase in rhodopsin-positive vesicles inside the RPE, while in the wild-type, rhodopsin antibodies only label continuous rod outer segments. Quantification confirmed that the lbk RPE contains increased numbers of outer segment-containing vesicles (Fig. 2T). Staining of eye cryosections with Oil Red O confirmed the presence of numerous lipid-containing vesicles in the lbk RPE (Fig. 4J). These data further suggest a failure in the mutant RPE to metabolise phagocyted PRC membrane stacks.

At 5 dpf, the PRC outer segments are fully developed in the zebrafish. In lbk mutants, however, they are reduced in length and their regular palisade arrangement is disrupted (Fig. 2I-L,U). Moreover, the microvilli from the RPE that normally interdigitate with the outer segments, cannot be detected. As development proceeds, these phenotypes get stronger resulting in PRCs with very short or even no outer segments (Fig. 2L).

Although we found the outer retina to be severely affected, the inner retina showed no evidence of morphological alterations in lbk (Fig. 2A-F). Similarly, staining of retinal Müller glia cells, which span the entire inner retina and are thus a good marker for inner retinal architecture, revealed no differences in their number, arrangement and morphology between wild-type and lbk<sup>−/−</sup> larvae (see Fig. S6 in the supplementary material). Starting at 7dpf, however, apoptotic cells can be detected in the neural retina (Fig. 2W).

To analyse the physiological effect of the lbk mutation on visual performance, we tested larvae by optokinetic response (OKR) measurements. In this test, eye tracking movements of immobilised larvae exposed to moving visual stimuli are used to assess visual ability. In contrast to wild-type larvae, moving gratings consistently failed to evoke eye movements in all lbk<sup>−/−</sup> larvae tested, indicating that mutants are behaviourally blind (Fig. 3A). However, this effect is not due to the inability of the mutants to move their eyes, as spontaneous eye movements were regularly observed. We further performed electroretinogram (ERG) analyses of 5 dpf wild-type and lbk larvae (Fig. 3B). In wild-type larvae the b-wave amplitude increased with increasing light intensities. In lbk, however, it remained small even at high light intensities. At 7dpf, the ERG of lbk<sup>−/−</sup> is flat, indicating that at this stage the mutant retina no longer responds to light stimuli (not shown).

**lbk larvae display vesicle phenotypes in the liver and intestinal tract**

Subsequent analyses revealed additional phenotypes in internal organs, including the liver and intestine. Starting at 6dpf, the liver of lbk larvae becomes progressively enlarged and darkly discoloured (Fig. 1J). Histological analyses confirmed this enlargement and showed a significant swelling of liver cells (Fig. 4A,B). This phenotype was confirmed when lbk<sup>−/−</sup> mutants were crossed to a transgenic line, Tg (ef1:GFP), that expresses GFP under the control of the intestinal promoter ef1 (Field et al., 2003) and sections imaged by confocal microscopy (Fig. 4C). To quantify the increase in liver size observed in lbk, we related the total liver area to the number of hepatocyte nuclei in a given section. By this measure, 5 dpf lbk larvae show a 30% increase of liver cell area compared to their siblings (Fig. 4D).

TEM analyses revealed that the cytoplasm of hepatocytes in lbk is filled with numerous, sometimes very large vesicles (up to 30 μm in diameter; Fig. 4E,F), indicative of a defect in vesicle trafficking in this cell type. At 7 dpf, the hepatocytes of lbk larvae show substantial necrotic changes (Fig. 4G,H). These changes might also explain the discolouration of the liver observable in external views (Fig. 1J). Oil Red O staining further showed an accumulation of vesicles containing large amounts of lipids in the lbk liver (Fig. 4I).

We further examined the intestines of lbk and age-matched sibling larvae. TEM analyses of 7 dpf larvae revealed a vesicle phenotype also in the cells of the mutant intestinal bulb. Although wild-type cells contain few large vesicles, the number of vesicles is significantly increased in lbk with a concomitant reduction in vesicle size (Fig. 4K,L; see Fig. S7 in the supplementary material). This further indicates a possible defect in the fusion of intracellular vesicles in lbk.

**The innate immune system is compromised in lbk mutants**

Similar to RPE cells, macrophages are highly phagocytotically active and contain numerous lysosomes and lysosome-related organelles, such as major histocompatibility complex class II compartments (Raposo and Marks, 2002). This prompted us to examine the morphology of vesicles in this cell type by incubating 7 dpf lbk and sibling larvae in Neutral Red, which is selectively retained in macrophage lysosomes (Herbomel et al., 2001). This staining suggests that intracellular vesicles in lbk macrophages may be enlarged, seem to display an amorphous shape and are more heterogeneous in size than in wild-type larvae (Fig. 5A,B; inserts).
Fig. 2. *lbk* larvae display a severely compromised RPE and shorter PRC outer segments. (A-H) Sections of *lbk* (B,D,F,H) and sibling (A,C,E,G) eyes show that overall eye development proceeds normally in *lbk*, including the formation of a multi-layered neural retina. Beginning at 4 dpf (A,D), however, the retinal pigment epithelium (RPE) is severely hypopigmented and becomes progressively inflated. At 7 dpf (E-H), the RPE in *lbk* is threefold thicker than in age-matched siblings (G,H; brackets). The inner retina in *lbk* is morphologically indistinguishable from wild-type siblings. INL, inner nuclear layer; PRC, photoreceptor cells; RGC, retinal ganglion cells. (I-N) TEM sections reveal that the *lbk* RPE contains very few mature melanosomes. The majority of melanosomes are aberrantly shaped and show regions that lack pigment. Moreover, the mutant RPE is filled with numerous vesicles. The number of vesicles increases from 5-7 dpf (compare J with L). The microvilli of the RPE (I, arrowheads) that normally interdigitate with the outer segments of PRCs are significantly reduced in *lbk* (J; arrow). Furthermore, the outer segment length is reduced. Some *lbk* PRCs virtually lack outer segments (L, arrowheads). (M,N) Higher magnification of vesicles containing undigested PRC outer segments (arrows) in the *lbk* RPE (N shows a higher magnification of the boxed area in M). M, PRC mitochondria; N, PRC nucleus; OS, PRC outer segment. (O,P) Merged phase contrast and fluorescence images of eye sections stained for rhodopsin (red) show an increase of discrete rhodopsin-positive vesicles in the *lbk* RPE (P, arrows) versus continuous PRC OS in the wild type (O, arrows). Insets show higher magnifications of parts of the RPE from the sibling and mutant retinas shown in O and P. (Q) Images showing melanosomes at different stages of maturation (I-IV) in the sibling (top) and *lbk* (bottom) RPE at 5 dpf. Arrowheads indicate melanosomes of the respective stages. (R-U) Statistical analyses of RPE thickness (R), average RPE cell area per nucleus (S), average number of vesicles containing PRC outer segments (T) and PRC outer segment length (U) in sibling and *lbk* larvae at 5 dpf. Error bars indicate standard deviations; *P* values were calculated using Student’s *t*-test. (V) Quantification of melanosomes at different stages of maturation in the *lbk* and sibling RPE at 5 dpf. Counts represent the numbers of melanosomes in areas of 10 μm². Although there are no substantial differences between the numbers of stage I-III melanosomes, mutant larvae contain significantly fewer mature (stage IV) melanosomes. (W) TUNEL labelling of cryosections shows increased levels of apoptosis in *lbk* (bottom right). The relatively dark appearance of the mutant RPE results from the thickness of these sections (40 μm). All images were derived from mid-transversal sections of the eye. Scale bars: 50 μm in A-F; 10 μm in G,H; 5 μm in I-M; 2.5 μm in N; 25 μm in O,P.

**lbk** resembles multi-systemic disorders
To obtain functional information on the innate immune system in *lbk*, we injected DsRed-expressing bacteria into mutant and sibling larvae at 48 hpf (Fig. 5C). Phagocytosis of the injected bacteria was complete within 10 minutes of the injection. Clearance of the fluorescent bacteria was monitored by in vivo fluorescence microscopy as well as by plating dissociated larvae onto selective media plates and counting of the resulting bacterial colonies. We found that while siblings efficiently cleared injected bacteria within 32 hours, mutants still contained numerous bacteria (Fig. 5D-M). Importantly, although phagocytosis appeared unaffected in *lbk*, the subsequent failure of the mutant phagocytes to degrade the bacteria indicates a defect in the fusion of phagosomes with lysosomes.

**Rescue and morpholino phenocopy of the *lbk* phenotype**

To confirm that a mutation in *vam6* underlies the *lbk* phenotype, we performed a rescue experiment. The zebrafish *vam6* cds was cloned into an expression vector under the control of a heatshock-inducible promoter (Bajoghli et al., 2004). This vector was injected at the one-cell stage and expression induced by heat shock at 3 dpf. In total, three independent experiments were performed and evaluated. To assess the rescue of the mutant phenotype, the larvae were screened at 5 dpf for their external morphology. *vam6*-injected *lbk* larvae developed darker skin melanocytes and RPE (Fig. 6A-C; insets) than un.injected and control vector-injected (no *vam6* cds) *lbk* larvae. The larvae were further tested by ERG analysis to assess the rescue of the visual ability. *vam6*-injected *lbk* larvae showed an increase in the b-wave amplitude to 20-60% of wild-type levels (Fig. 6A-D). By contrast, un injected control *lbk* larvae did not show a b-wave (Fig. 3B).

To provide further evidence that a mutation in *vam6* underlies the *lbk* phenotype, we designed an antisense morpholino against a region containing the start codon (ATG) of the *vam6* mRNA. Injection of the ATG-morpholino into wild-type embryos at the one-cell stage resulted in a similar, but stronger phenotype than that observed in *lbk* (Fig. 6E). Injection of a control morpholino did not result in larvae displaying any phenotypes (Fig. 6E). The weaker phenotype of *lbk* larvae compared with the *vam6* knock-down could be explained by the presence of maternally supplied wild-type *vam6* mRNA, which could serve as a template for the synthesis of functional Vam6p during early embryonic development. To test this hypothesis, we isolated mRNA from wild-type zygotes, reverse transcribed it into cDNA and performed a PCR analysis. This showed that *vam6* mRNA is maternally supplied in zebrafish (Fig. 6F) and its expression persists through later stages of development (Fig. 6F,G; see Fig. S8 in the supplementary material). The Vam6p protein is part of the HOPS complex, a large multi-subunit protein complex. In this context it is interesting that the mRNA for Vps18, another member of the HOPS complex, has been found to be maternally supplied in zebrafish zygotes in a microarray experiment (http://zf-espresso.tuebingen.mpg.de/).

**DISCUSSION**

In this study, we provide the first phenotypic description of a lack of *vam6* gene function in a multicellular organism. Zebrafish *lbk* mutants display defects in several tissues and cell types (liver, intestinal tract, macrophages, PRCs, RPE and skin melanocytes) as well as compromised vision and innate immunity. The range of phenotypes resembles pathological symptoms observed in individuals suffering from syndromes caused by defects in lysosomes and lysosome-related organelles (Stinchcombe et al., 2004), including the ARC syndrome, CHS, GS and HPS. For example, CHS, GS and HPS are characterised by a hypopigmentation of the skin and RPE, as well as a significant visual impairment (Barak and Nir, 1987; Clark and Griffiths, 2003; Griscelli et al., 1978; Hermansky et al., 1975). These symptoms are also observed in *lbk* mutants.

Melanosomes are part of the large group of lysosome-related organelles and serve to synthesize and store melanin. The melanocytes in *lbk* mutants display significantly fewer and lighter melanosomes than are observed in wild-type animals. The presence of melanosomes with wild-type levels of melanin pigment in...
the lbk skin and RPE shows that the mutation does not cause a block in melanin synthesis or melanosome biogenesis per se. The increase in immature melanosomes and the substantially decreased number of mature (stage IV) melanosomes in lbk rather suggests a significantly reduced efficiency of melanosome maturation, a process critically dependent on the fusion of endosomal vesicles with premelanosomes (Dell’Angelica, 2003). Alternatively, the stop codon mutation we identified in lbk, UAG, has been suggested to have intermediate read-through fidelity. Thus, the mutant embryos might retain a low level of functional protein, as has been observed in the human syndromes described in this manuscript and other lysosomal storage disorders (Brooks et al., 2006).

The hypopigmentation of the lbk RPE probably contributes to the reduced visual ability. Moreover, the RPE contains numerous vesicles containing undigested PRC outer segments. Although indicating that phagocytosis by the RPE is not compromised, this novel phenotype suggests a defect in lysosome-mediated degradation of phagocytosed outer segments, a central function of the RPE, which may lead to a generalised metabolic dysregulation. As the RPE is essential for maintaining the PRCs, compromising the viability of the RPE will probably also negatively affect PRC function. This notion is supported by our TEM studies, which reveal shortened PRC outer segments. The presence of an a-wave in ERGs, however, suggests that lbk PRCs retain at least some functionality.

Fig. 4. lbk larvae display vesicular phenotypes in liver and intestinal cells. (A,B) Longitudinal sections showing the liver and part of the intestine. There is a substantial increase in liver size (outlined) in lbk. Sections were derived from the same region of the body and images are reproduced at the same magnification. (C) Confocal LSM images of larvae expressing GFP under the intestinal promoter ef1 reveal an increase in liver size (top row, outlined) in lbk. Nuclei are stained with DAPI (blue). Bottom row shows higher magnifications of regions from livers shown in the top row. Individual liver cells are outlined showing the increase in hepatocyte size in lbk. The arrangement of hepatocytes in lbk is disorganised. (D) Quantification of the average cell area per nucleus in histological sections of livers revealed that the lbk liver shows on average an area 30% larger than that of age-matched siblings [n=5 (sibs.), n=7 (lbk); P=0.0001]. (E-H) Transverse TEM sections of livers show that lbk hepatocytes contain numerous large vesicles (v). Asterisks indicate erythrocytes. At 7 dpf, lbk hepatocytes display necrotic changes: e.g. a condensation of nuclear chromatin; an increase in the space between the nuclear membranes (arrowheads); a massively inflated ER lumen (#), which is continuous with the nuclear membranes; and numerous vesicles often displaying an electron-dense lumen. N, nucleus. (I,J) Oil Red O staining of the liver (I) and RPE (J) reveals lipid-containing vesicles (arrowheads) in hepatocytes and the RPE in lbk. (K,L) Transverse TEM sections of the anterior intestine show an increase in number and decrease in size of vesicles in the cells lining the intestine in lbk. Scale bars: 50 μm in A,B; 50 μm in C; 25 μm in insets; 10 μm in E,F,I,J; 5 μm in G,H; 2 μm in K,L.
Fig. 5. *lbk* larvae show a compromised ability to clear bacterial infections. (A,B) Dorsal views of larvae in which macrophages were stained with Neutral Red to reveal intracellular vesicles (arrows). Compared with the vesicles observed in wild-type macrophages, those in *lbk* appear enlarged, heterogeneous in size and less regularly shaped. Insets show higher magnifications of macrophage vesicles. (C) Combined bright-field and fluorescence images of the tail of a 48 hpf larva 5 minutes after the injection of DsRed-fluorescent bacteria, showing bacteria still circulating in the bloodstream (inset; fluorescent traces of bacteria moving in blood vessels) and bacteria that have been engulfed by phagocytes (arrowheads). (D-G) Head regions of sibling and *lbk* larvae 32 hours after bacteria injection. Fluorescent bacteria-laden phagocytes are present in *lbk* (arrowheads). (H-J) Bright-field (H) and fluorescence images (I) of a bacterial plate showing bacterial colonies from an *lbk* larva macerated at 85 hpf, 37 hours after *Salmonella* injection. (J-M) Tails of sibling and *lbk* larvae 32 hours after bacteria injection. Arrowheads indicate fluorescent bacteria-containing phagocytes.

up until 5 dpf. Nonetheless, *lbk* larvae are severely visually impaired. Visual defects have also been reported for CHS and HPS syndromes (OMIM #214500; #203300). Interestingly, visual problems have not been diagnosed in individuals with GS to date (OMIM #214450; #607624; #609227). This may, however, be due to inconsistent testing of affected individuals or subsymptomatic reductions in visual acuity.

In addition to the RPE, there is also an accumulation of vesicles in hepatocytes and the cells lining the intestinal tract. These cells typically show high endo- and exocytic activities and large numbers of intracellular lysosomes and lysosome-related vesicles. The larger number of intracellular vesicles in the *lbk* liver and RPE is accompanied by an increase in overall liver and RPE size. As there is no apparent defect in the formation of vesicles but rather in their subsequent fusion (and hence recycling), it could be envisaged that the observed increase is due to an accumulation of vesicles, which in turn leads to an overall increase in cytoplasmic volume. An increase in liver volume (hepatomegaly) has also been described for individuals affected by CHS (OMIM #214500) and ARC syndrome (Gissen et al., 2004). Similarly, the intestinal phenotype observed in *lbk* is reminiscent of the granulomatous colitis described for individuals with HPS (OMIM #203300).

Defects in lysosome-related organelles might also be responsible for the immunological deficiencies observed in individuals with CHS and GS (Faigle et al., 1998; Griscelli et al., 1978). Macrophages, for instance, are crucial for the body’s innate immune response. One of their key functions is the phagocytosis of pathogens. Pathogen-containing endosomes subsequently fuse with lysosomes and major histocompatibility complex class II (MHCII) compartments, a lysosome-related organelle, in which the pathogens are degraded and foreign peptides loaded onto MHCII complexes. Loaded MHCII complexes are subsequently delivered to the surface of macrophages for presentation to cells of the adaptive immune response. We found that *lbk* macrophages display an increase in intracellular vesicles, which might be indicative of a defect in the endosomal pathway and/or a defect in MHCII vesicle delivery to the plasma membrane. Importantly, our analyses show that the innate immune response to bacterial infection is significantly compromised in *lbk*.

ARC syndrome, CHS, GS and HPS are caused by defects in vesicle trafficking, particularly of lysosomes, lysosome-related organelles and late endosomes, and mutations in genes involved in vesicle transport, sorting, docking and fusion were shown to underlie these diseases. Moreover, ARC syndrome was demonstrated to be caused by mutations in the *vps33b* gene, a member of the HOPS complex, in humans (Gissen et al., 2004). Similarly, morpholino knock-down experiments targeting the zebrafish *vps33b* gene result in a phenotype resembling ARC syndrome (Matthews et al., 2005). Interestingly, the buff (*bf*) mouse, which harbours a mutation in the *Vps33a* gene, has been proposed as a model for HPS (Suzuki et al., 2003), suggesting a link between ARC syndrome and the CHS, GS and HPS group of diseases. The phenotypes observed in *lbk* encompass both symptoms observed in individuals with CHS, GS and HPS group of diseases. Moreover, we have identified the gene mutated in *lbk* as *vam6*. Like Vps33p, the Vam6p protein is a component of the HOPS complex, which is required for SNARE complex assembly during vesicle docking and fusion. Mutations in the two alleles identified in this study lead to a premature STOP upstream of the CLH repeat domain and Ypt7p-interaction sequences, and the exchange of a highly conserved amino acid in the N-terminal CNH domain, respectively. The latter has been shown to be essential for lysosome clustering and fusion (Caplan et al., 2001). The Ypt7p-interaction domain mediates protein-protein interactions between Vam6p/Vps39p and Ypt7p/Rab7, a Rab-GTPase with essential functions in early to late endosome transport and late endosome-
Fig. 6. The lbk phenotype can be partially rescued by the expression of vam6 in lbk–/– larvae and phenocopied by vam6 knock-down. (A-C) Electrophoretogram analyses of 5 dpf vam6 vector-injected and heat-shocked sibling (A) and lbk–/– (B, C) larvae. In contrast to uninjected lbk larvae (fig. 3B), injected lbk larvae displayed significant b-waves, indicating partial rescue of retinal function. The b-wave amplitude varied between different lbk larvae with some showing up to 60% of the wild-type b-wave amplitude (B), while others reach only ~20% (C). Insets show head regions of vam6 vector-injected sibling (A) and lbk larvae (B, C) and the extent of rescue in RPE and skin melanoocytes pigmentation. Arrowheads indicate skin melanoocytes with near wild-type levels of melanin in rescued larvae. (D) Quantification of the observed ERG rescue: sibling larvae display an average b-wave amplitude of 286 μV; vam6 vector-injected lbk larvae show an average b-wave amplitude of 83 μV (n = 7 for both sets of larvae; error bars show standard deviations). (E) Phenotype of the vam6 knock-down at 36 hpf showing the hypopigmentation of the RPE and skin melanoocytes (insets; arrowheads) characteristic of lbk. However, the knock-down results in additional phenotypes not observed in lbk, including a small head and eyes and a shortened body axis. Larvae displaying wild-type levels of melanin are age-matched control morpholino-injected individuals. (F) PCR analysis for the presence of vam6 transcripts in RNA isolated from wild-type zebrafish zygotes (upper half of gel) and 24 hpf embryos (lower half). β-Actin (expressed at all developmental stages) and pmel17 onset of expression (~20 hpf [Schonthaler et al., 2005]) served as controls. Lane M, 100 bp ladder; lane 1, amplification of a 530 bp β-actin cDNA fragment; lanes 2 and 4, amplification of two different fragments of the zebrafish vam6 cDNA (2628 bp and 707 bp) with independent primer pairs; lane 3, amplification of a 2538 bp fragment of the pmel17 cDNA. (G) Whole-mount in situ hybridisation on PTU-treated 48 hpf wild-type embryos (left, lateral view; right, dorsal view) showing vam6 expression.

lysosome fusion, as well as in axonal retrograde transport (Deinhardt et al., 2006; Stein et al., 2003). It has been shown in yeast that Vam6p acts as a guanine nucleotide exchange factor for Ypt7p (Wurmser et al., 2000). When this exchange fails, Ypt7p is locked in its GDP-bound state and vesicle docking is blocked. This probably contributes to the vesicle phenotypes observed in lbk. In mammalian cells, both Rab7 and the HOPS complex are essential for the conversion of early into late endosomes (Rink et al., 2005). Interestingly, RNAi-mediated knock-down of Rab7 in HeLa cells leads to a cellular vesicle phenotype reminiscent of that observed in fibroblasts from individuals with CHS (Davies et al., 1997). Similarly, knock-down of Rab7 in C. elegans leads to enlarged early and late endosomes and knock-down of HOPS complex members yield defects in gut lysosome formation (Poteryaev et al., 2007).

The next step in vesicle fusion, following the initial docking steps, is assembly of SNARE complexes. This process requires physical interactions between the HOPS complex and SNAREs (Collins et al., 2005; Price et al., 2000a; Price et al., 2000b; Sato et al., 2000). Mutations in Vam6p that abolish the functions of the CLH or CNH domains, respectively, might hamper these interactions and additionally contribute to the accumulation of vesicles in lbk.

The mutation in the vam6 gene in lbk acts in a cell-autonomous fashion and affects cells derived from all three germ layers (skin melanocytes, RPE, PRCs in ectoderm; liver and intestine in endoderm; and macrophages in mesoderm). The most severe phenotypes are detected in cells that display high levels of phagocytic (RPE cells, macrophages) and secretory activity (hepatocytes, intestinal cells). Owing to the importance of vesicle trafficking in these cells, they are probably the first and most severely affected. The occurrence of only partially overlapping symptoms in ARC syndrome, CHS, GS and HPS, as well as the zebrafish lbk mutant could be due to the fact that lysosome-related organelles can be cell type-specific (Rapoport and Marks, 2002) and thus defects in proteins required for these organelles might only affect certain cell types. The comparatively widespread organ dysfunction observed in lbk suggests that Vam6p is a core component with non-redundant functions in vesicle trafficking in a range of cell types.

It has recently been shown that a viral insertion disrupting the vps18p gene, another member of the HOPS complex, in zebrafish results in similar phenotypes as those observed in lbk, including a hypopigmentation of skin and RPE melanocytes, a lack of iridophore reflections, visual defects, hepatomegaly and an accumulation of vesicles in hepatocytes (Maldonado et al., 2006; Sadler et al., 2005). This provides further evidence for the importance of the HOPS complex in various cell types and organs.

In conclusion, the overlap in symptoms in human syndromes caused by defective lysosome trafficking and phenotypes observed in lbk suggest this mutant as a model for these inherited human
diseases and suggests a novel disease gene. Importantly, there is currently no mouse model for loss of function of Vam6p. Studies on lbk may offer important insights into the mechanisms underlying the observed symptoms and make these pathologies amenable to experimental manipulation.

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Supplementary material
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