SLC26A4 Targeted to the Endolymphatic Sac Rescues Hearing and Balance in Slc26a4 Mutant Mice

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Abstract

Mutations of SLC26A4 are a common cause of human hearing loss associated with enlargement of the vestibular aqueduct. SLC26A4 encodes pendrin, an anion exchanger expressed in a variety of epithelial cells in the cochlea, the vestibular labyrinth and the endolymphatic sac. Slc26a4−/− mice are devoid of pendrin and develop a severe enlargement of the membranous labyrinth, fail to acquire hearing and balance, and thereby provide a model for the human phenotype. Here, we generated a transgenic mouse line that expresses human SLC26A4 controlled by the promoter of ATP6V1B1. Crossing this transgene into the Slc26a4−/− line restored protein expression of pendrin in the endolymphatic sac without inducing detectable expression in the cochlea or the vestibular sensory organs. The transgene prevented abnormal enlargement of the membranous labyrinth, restored a normal endocochlear potential, normal pH gradients between endolymph and perilymph in the cochlea, normal otocochlea formation in the vestibular labyrinth and normal sensory functions of hearing and balance. Our study demonstrates that restoration of pendrin to the endolymphatic sac is sufficient to restore normal inner ear function. This finding in conjunction with our previous report that pendrin expression is required for embryonic development but not for the maintenance of hearing opens the prospect that a spatially and temporally limited therapy will restore normal hearing in human patients carrying a variety of mutations of SLC26A4.

Introduction

Enlargement of the vestibular aqueduct (EVA; OMIM #600791) is a malformation of the temporal bone that is commonly observed in children with sensorineural hearing loss [1,2,3,4,5]. Mutations of SLC26A4 are the most common cause for EVA-associated hearing loss that can either be non-syndromic (DFNB4; OMIM #600791) or syndromic with enlargement of the thyroid gland (Pendred syndrome; OMIM #274600). SLC26A4 codes for the anion exchanger pendrin that transports anions such as Cl−, I− and HCO3− [6,7]. Although EVA is a malformation of the temporal bone, it is not the cause for hearing loss since no correlation was found between the degree of EVA and the severity of hearing impairment [8]. EVA, however, is an indication of an enlargement of the endolymphatic duct epithelium that was present during embryonic development. Cartilage cells that form in the periphery of the endolymphatic duct epithelium preserve the diameter of the duct in a ‘fossil-like’ record when they give rise to the bone of the vestibular aqueduct.

The mature inner ear consists of seven interconnected fluid spaces that house six sensory organs (Fig. 1): The cochlea for hearing, the utricle and saccule for sensing linear acceleration including gravity, and three ampullae with semicircular canals for sensing angular acceleration in three spatial axes. The seventh fluid compartment is the endolymphatic duct and sac, which is devoid of sensory cells and which is suspected to play a role in fluid homeostasis [9,10]. Pendrin is expressed in a variety of epithelial cells that enclose endolymph, which is the luminal fluid of the inner ear (Fig. 1). Pendrin is expressed in outer sulcus, spiral prominence and spindle-shaped cells in the cochlea, transitional cells in the utricle, saccule and ampullae and mitochondria-rich cells (synonym: Forkhead-related or FORE cells) of the endolymphatic sac.
Mutations of SLC26A4 are the most common cause for hearing loss associated with a swelling of the inner ear. This human disease is largely recapitulated in a mutant mouse model. Mutant mice lack Slc26a4 expression and their inner ears swell during embryonic development, which leads to failure of the cochlea and the vestibular organs resulting in deafness and loss of balance. SLC26A4 is normally found in the cochlea and vestibular organs of the inner ear as well as in the endolymphatic sac, which is a non-sensory part of the inner ear. The multitude of sites where SLC26A4 is located made the goal to restore function through restoration look futile, unless some sites were more important than others. Here, we generated a new mutant mouse that expresses SLC26A4 in the endolymphatic sac but not in the cochlea or the vestibular organs of the inner ear. Fantastically, this mouse did not develop the detrimental swelling of the inner ear and even more exciting, the mouse developed normal hearing and balance. Our study provides the proof-of-concept that a therapy aimed at repairing the endolymphatic sac during embryonic development is sufficient to restore a life-time of normal hearing and balance.

Figure 1. Schematic diagram of the inner ear. A) Diagram of the membranous labyrinth. The two continuous luminal fluid spaces of the mature inner ear are filled with endolymph (pink and purple). B–E) Diagrams of a cross section of one cochlear turn (B), of the utricle or saccule (C), of one ampulla (D) and the endolymphatic sac (E). Cells that express pendrin (yellow cells pointed to by arrows) are diagrammed in mature tissues.

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the Slc26a4+/A line to generate mice that lack expression of mouse pendrin but express human pendrin in the endolymphatic sac. No expression of pendrin protein was detected in these mice in the cochlea or the vestibular labyrinth but in mitochondria-rich cells of the endolymphatic sac. Analysis of this mouse model revealed normal hearing and balance function. Our data indicate that the expression of pendrin solely in the endolymphatic sac of the inner ear is sufficient to permit the development of normal hearing and balance.

Results

Generation of Tg(+);Slc26a4+/A transgenic mice

A transgenic mouse line, referred to as Tg(B1-hPDS)1x+/; Slc26a4+/A and abbreviated here to Tg(+);Slc26a4+/A, was created by the laboratory of Dr. Dominique Eladari (Paris, France) [15]. This mouse expresses human SLC26A4 (formerly named hPDS) controlled by the promoter of ATP6V1B1, which codes for the B1-subunit of the vH-ATPase. Transgenic founders were crossed with wild-type C57BL/6 x CBA F1 mice and three Tg(+);Slc26a4+/A mice were shipped to Kansas State University (Manhattan, Kansas, USA). At Kansas State University, Tg(+);Slc26a4+/A mice were crossed with Slc26a4+/A mice, which are maintained in an isogenic 129S6SvEv background, to generate the desired Tg(+);Slc26a4+/A mice in an F2 generation. Expression of SLC26A4 (human pendrin) in this mouse was expected to originate solely from the transgene since Exon 8 in the Slc26a4 allele was replaced with a neomycin-cassette that introduced a frame-shift [14]. Littermates with the genotype Tg(−);Slc26a4+/A served as negative controls. These mice were expected to lack functional pendrin protein expression. Further, littermates with genotypes Tg(−);Slc26a4+/A, Tg(+);Slc26a4+/A, and Tg(+);Slc26a4+/A served as positive controls. These mice were expected to express murine pendrin with or without augmentation of human pendrin and have normal hearing and balance.

mRNA expression

Expression of Atp6v1b1, Atp6v1b1, Slc26a4 and SLC26A4 was determined by quantitative RT-PCR and normalized to the expression of 18S rRNA (Fig. 2). The highest levels of Atp6v1b1 and Slc26a4 mRNA among the different inner ear tissues were found in the endolymphatic sac (Fig. 2B and C). Expression of Slc26a4 was reduced by factors between 6 and 16 in Tg(+);Slc26a4+/A mice compared to Tg(−);Slc26a4+/A mice (Fig. 2C vs G). Expression levels of Atp6v1a1 and Atp6v1b1 exhibited a similar pattern among inner ear tissues of Tg(−);Slc26a4+/A and Tg(+);Slc26a4+/A mice (Fig. 2A vs E and Fig. 2B vs F). Most interesting, expression levels of human SLC26A4 in Tg(+)Slc26a4+/A resembled the pattern of mouse Slc26a4 in Tg(−);Slc26a4+/A mice with the highest levels being expressed in the endolymphatic sac (Fig. 2C vs H). Whether or not expression levels of human SLC26A4 in Tg(+);Slc26a4+/A exceeded expression levels of mouse Slc26a4 in Tg(−);Slc26a4+/A mice remained undetermined, since the efficiency of the reverse transcription of

Figure 2. Atp1a1, Atp6v1b1, Slc26a4, and SLC26A4 mRNA levels in inner ear tissues. Expression was determined by quantitative RT-PCR performed on total RNA. Total RNA was isolated from microdissected tissues obtained from Tg(+/+);Slc26a4+/A (A–D) and Tg(+/+);Slc26a4+/A mice (E–H). Endolymphatic sacs (ES) were isolated from mice at age E17.5. Cochleae were isolated at ages E17.5 (C1) and P2 (C2). Vestibular labyrinths (VL), consisted of saccule, utricle, ampulla and semicircular canals without endolymphatic sacs, were isolated at age P8. The expression of endogenous mouse Atp1a1, Atp6v1b1, and Slc26a4, and of transgenic human SLC26A4 mRNA was normalized to the expression of 18S rRNA. Note that the expression pattern of human SLC26A4 in Tg(+/+);Slc26a4+/A mice resembles the pattern of mouse Slc26a4 in Tg(−);Slc26a4+/A mice (both patterns highlighted in red). Numbers inside graphs represent the number of experiments.

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mRNA into cDNA remains generally unknown in quantitative RT-PCR experiments. Taken together, the data demonstrate that the transgene restored pendrin mRNA expression to the endolymphatic sac, the cochlea and the vestibular labyrinth of the inner ear.

**Protein expression - eGFP expression**

The ability of the *ATP6V1B1* promoter to drive protein expression in different tissues including the cochlea, the vestibular labyrinth and the endolymphatic sac was evaluated in a transgenic mouse line, Tg(*B1-eGFP*) in which the expression of eGFP is controlled by the same 6.9 kb promoter of the human *ATP6V1B1* gene that drives the expression of human pendrin in Tg(+);Slc26a4Δ/Δ mice [16]. No expression of eGFP was detected in the cochlea or the vestibular labyrinth of E15.5 Tg(*B1-eGFP*) mice, although expression was present in the endolymphatic sac and the kidney (Fig. 3). These data suggest that the *ATP6V1B1* promoter does not drive protein expression in the cochlea or the vestibular labyrinth.

**Protein expression - Western blotting**

Soft tissues of the cochlea and the vestibular labyrinth, exclusive of the endolymph sac, were collected from adult mice by microdissection and pooled into an ‘inner ear’ sample. Crude membrane protein preparations were obtained from these inner ear samples and from kidneys and subjected to gel-electrophoresis and Western blotting. Membrane proteins were obtained from Tg(+);Slc26a4Δ/Δ mice as well as from Tg(−);Slc26a4Δ/Δ mice, which served as positive controls, and from Tg(−);Slc26a4Δ/Δ mice, which served as negative controls. Pendrin was detected in the inner ear and kidney of Tg(−);Slc26a4Δ/Δ mice as a ~110 kDa band (Fig. 4A). Inner ear from Tg(+);Slc26a4Δ/Δ mice lacked this band. The observation that there was no difference in the pattern of faint bands between inner ears from Tg(+);Slc26a4Δ/Δ mice and Tg(−);Slc26a4Δ/Δ mice, which is the negative control, suggests that pendrin was either not detectable or not present. The pendrin band, however, was found in kidney from Tg(+);Slc26a4Δ/Δ mice (Fig. 4B), which suggests that the antibody recognizes both mouse and human pendrin. The observation that pendrin was detected at

![Figure 3. Transgene-encoded eGFP expression. A–C Expression of eGFP was monitored as direct fluorescence in cochlear ducts (A), endolymphatic sacs (B), and kidney slices (C) freshly isolated from E15.5 Tg(*B1-eGFP*) mice. The outline of the imaged tissues is marked (dashed-line). D–I Expression of eGFP was evaluated by immunocytochemistry in the cochlea (D), utricle (E), ampulla (F), endolymphatic sac (G), kidney cortex (H), and kidney medulla (I) from E15.5 Tg(*B1-eGFP*) mice. Staining consisted of immunocytochemistry of eGFP (green), F-actin (red) and nucleic acids (blue). The number of mice represented by these images are 2 for images A–C and 2 for images D–I. MR, mitochondria-rich cells; IC, intercalated cells; Lim, spiral limbus; K, Kolliker’s organ; OS, outer sulcus; SV, stria vascularis; HC, vestibular hair cells; TC, transitional cells; M, melanocytes. doi:10.1371/journal.pgen.1003641.g003](image-url)
similar levels in descending amounts of kidney proteins isolated from Tg(−);Slc26a4−/+ mice (Fig. 4A) and Tg(+);Slc26a4+/A mice (Fig. 4C) suggests that the detection threshold for mouse and human pendrin was similar. Whether the antibody differed in the sensitivity between mouse and human pendrin remains unknown, since the relative abundance of mouse pendrin in kidneys of Tg(−);Slc26a4+/D mice and human pendrin in kidneys of Tg(+);Slc26a4+/A mice is not known.

The meaning of pendrin being not detectable in the inner ear was evaluated by comparison of the intensity of the pendrin band in inner ear to the intensities in descending amounts of kidney protein (Fig. 4A). This comparison suggests that a 5-fold lower amount pendrin should have been detectable in the inner ear. This means that pendrin expression in the inner ear of Tg(+);Slc26a4+/A mice is either absent or expressed at a level that does not exceed 20% of the expression level in Tg(−);Slc26a4−/+ mice.

Paint-fill of the cochlea and the endolymphatic sac

Temporal bones from Tg(+);Slc26a4+/A and Tg(+);Slc26a4+/D mice were isolated at E15.5, fixed and injected with white paint (Fig. 5A and B). Most striking is that there was no enlargement of the endolymphatic sac, duct or cochlea in Tg(+);Slc26a4+/A mice and that the morphology of Tg(+);Slc26a4+/A and Tg(+);Slc26a4+/D mice was grossly similar. These data demonstrate that the introduction of the transgene rescued the malformation previously described in Slc26a4+/A mice [10,14].

Figure 4. Pendrin expression evaluated by Western blotting. Proteins in crude membranes prepared from pooled inner ear tissues consisting of soft tissues from the cochlea and the vestibular labyrinth devoid of endolymphatic sac and crude membranes prepared from kidneys of Tg(−);Slc26a4−/+ , Tg(−);Slc26a4+/A and Tg(+);Slc26a4+/A mice were resolved by gel-electrophoresis and detected with an anti-pendrin antibody (Pds #1) and an anti-β-actin antibody. β-actin served as a loading control. A) Comparison of pendrin expression in membranes prepared from inner ears of Tg(−);Slc26a4−/+ , Tg(−);Slc26a4+/A and Tg(+);Slc26a4+/A mice to expression levels in descending amounts of membranes prepared from kidneys of Tg(−);Slc26a4−/+ mice. B) Comparison of pendrin expression in membranes prepared from kidneys of Tg(−);Slc26a4−/+ , Tg(−);Slc26a4+/A and Tg(+);Slc26a4+/A mice. C) Expression levels of pendrin in descending amounts of membranes prepared from kidneys of Tg(−);Slc26a4−/+ mice. Data shown in A, B and C are each representative of 2 biological replicates.

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Histology and pendrin expression in the endolymphatic sac

Whole-mounted specimens of the endolymphatic sac were prepared for immunocytochemistry from Tg(+);Slc26a4Δ/Δ, Tg(−);Slc26a4Δ/Δ, and Tg(+);Slc26a4Δ/+ mice (Fig. 5C–E). Most striking is the enlargement and lack of pendrin expression in the endolymphatic sac of Tg(−);Slc26a4Δ/Δ mice (Fig. 5E) and the similarity in size and similarity in pendrin expression between the endolymphatic sac of Tg(+);Slc26a4Δ/Δ mice (Fig. 5E) and Tg(+);Slc26a4Δ/+ mice (Fig. 5D). These data demonstrate that the transgene drives pendrin expression in the endolymphatic sac and that the introduction of the transgene rescued the malformation [10,14].

Gross morphology of the cochlea

Gross morphological examination of inner ears revealed greater similarity between Tg(+);Slc26a4Δ/Δ mice and Tg(+);Slc26a4Δ/+ than between Tg(+);Slc26a4Δ/Δ mice and Tg(−);Slc26a4Δ/Δ mice (Fig. 6A–C). Cochlear turns in Tg(+);Slc26a4Δ/Δ mice appeared normal in width and did not show widening of turns or thinning of the otic capsule that was seen in Tg(−);Slc26a4Δ/Δ and that was previously described in Slc26a4Δ/Δ mice [12,17]. Inspection of the oval window revealed ‘glittering’ otoconia in the saccule in Tg(+);Slc26a4Δ/Δ and Tg(+);Slc26a4Δ/+ mice in contrast to Tg(−);Slc26a4Δ/Δ mice where no ‘glittering’ was visible (Fig. 6D–F).

Histology and pendrin expression in the cochlea

Midmodiolar sections of cochlear tissues were prepared for immunocytochemistry from Tg(+);Slc26a4Δ/Δ mice and positive controls consisting of Tg(−);Slc26a4Δ/+ or Tg(+);Slc26a4Δ/+ mice. No evidence for cochlear enlargement was found in Tg(+);Slc26a4Δ/Δ mice at E16.5 (Fig. 7B), P1 (Fig. S1A), P16 (Fig. 7A; Fig. S1A), or P18 (Fig. S1C) suggesting that the introduction of the transgene rescued the cochlear malformation previously described in Slc26a4Δ/Δ mice, which includes a ∼10-fold enlargement of the cochlea [12,14]. No detectable pendrin expression was found in the spiral prominence or outer sulcus epithelium of the cochlea in Tg(+);Slc26a4Δ/Δ mice although prominent expression was observed in these cells in positive controls (Fig. 7 and S1). The absence of pendrin in Tg(+);Slc26a4Δ/Δ mice was observed with two different anti-pendrin antibodies (Pds #1 and Pds #2). The patterns of pendrin expression in the positive controls, Tg(−);Slc26a4Δ/+ and Tg(+);Slc26a4Δ/+ mice, were similar for both antibodies to the pattern previously observed in Slc26a4Δ/Δ mice [11,12,13]. Expression of pendrin was further examined in whole-mounted
Endocochlear potential and pH

The endocochlear potential and the difference in pH between endolymph and perilymph was measured with double-barreled ion selective electrodes in Tg(−);Slc26a4Δ/Δ mice, Tg(+);Slc26a4Δ/Δ mice, and Tg(+);Slc26a4Δ/Δ mice [18]. In contrast, Tg(−);Slc26a4Δ/Δ mice, as reported for Slc26a4Δ/Δ mice [18], developed a normal endocochlear potential and a normal endolymphatic pH that was higher (more alkaline) than in perilymph. Similar to Tg(+);Slc26a4Δ/Δ mice, Tg(−);Slc26a4Δ/Δ mice developed a normal endocochlear potential and a normal endolymphatic pH even though no detectable pendrin expression was observed in the cochlear epithelium. These data demonstrate that the introduction of the transgene, which rescued the malformation, also rescued the loss of the endocochlear potential and the loss of normal endolymphatic pH homeostasis.

Hearing

Hearing tests were based on auditory brain stem recordings and thresholds in response to tone bursts of 8 kHz, 16 kHz, and 32 kHz. Tests performed in Tg(+);Slc26a4Δ/Δ, Tg(−);Slc26a4Δ/Δ, and Tg(−);Slc26a4Δ/Δ mice confirmed profound deafness in Tg(−);Slc26a4Δ/Δ mice (Fig. 9B) consistent with previous findings in Slc26a4Δ/Δ mice [14, 18]. Waveforms of auditory brain stem recordings as well as thresholds were similar between Tg(+);Slc26a4Δ/Δ mice (Fig. 9A) and Tg(+);Slc26a4Δ/Δ (Fig. 9C). These findings demonstrate that the introduction of the transgene rescued normal hearing although the cochlea did not express detectable levels of pendrin. We next evaluated whether the rescued hearing phenotype in Tg(+);Slc26a4Δ/Δ would be stable through at least 3 months of age. Audiatory brain stem recordings were performed in Tg(+);Slc26a4Δ/Δ mice (Fig. 9D–F), in Tg(+);Slc26a4Δ/Δ mice (Fig. 9G–I), and in Tg(+);Slc26a4Δ/Δ mice (Fig. 9D–F) at 1, 2, and 3 month of age. Hearing in Tg(+);Slc26a4Δ/Δ mice at 8 kHz and 16 kHz was stable through 3 months.
Figure 7. Histology and pendrin expression in the cochlea. Staining in all images consisted of immunocytochemistry of pendrin (Pds #1 antibody; red), F-actin (green) and nucleic acids (blue). Images provide comparison of cryosections (A–B) and whole-mounted specimens (C–E) from Tg(+) Slc26a4Δ/Δ mice to cryosections (F–G) and whole-mounted specimens (H–J) from Tg(−) Slc26a4Δ/Δ mice. Whole-mounted specimens in C, E, H and J were imaged by detecting fluorescence in 25 optical sections that were recorded in 1 μm intervals and projected into a single plane. Whole-mounted specimens in D and I were imaged by detecting fluorescence in 8 optical sections that were recorded in 1 μm intervals, projected into a single plane.
single plane and overlaid onto a single brightfield image. The number of pairs of mice represented by these images are 2 for image A & F, 3 for B & G, and 1 for C–E & H–J with 3 sections being evaluated per animal. K, Kölliker’s organ; OS, outer sulcus; LS, limbus spiralis; IS, inner sulcus; OC, organ of Corti; SP, spiral prominence; SV, stria vascularis; RM, Reissner’s membrane. Additional images using an alternative anti-pendrin antibody (Pds2) and alternative positive controls (Tg(+);Slc26a4+/− mice) are provided in the Supplement (Fig. S1).

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Thresholds were very similar among individuals and did not differ from Tg(+);Slc26a4+/− and Tg(+);Slc26a4+/− mice. A greater variability in hearing thresholds was observed at 32 kHz (Fig. 9L), with 10 of the 19 Tg(+);Slc26a4+/− mice maintaining excellent hearing (thresholds ≤30 dB at 32 kHz) and 5 developing a high-frequency hearing loss (thresholds ≥60 dB at 32 kHz). About one half of the Tg(+);Slc26a4+/− mice (9 of 19) developed progressive threshold elevations at 32 kHz with thresholds increasing by ≥10 dB between the monthly measurements. This variability is reflected in the greater error bars at 32 kHz but did not lead to a statistically significant difference between Tg(+);Slc26a4+/− and Tg(+);Slc26a4+/− mice.

Histology and pendrin expression in the vestibular labyrinth

Sections and whole-mounted specimens of vestibular tissues were prepared for immunocytochemistry from Tg(+);Slc26a4+/− mice and positive controls consisting of Tg(−);Slc26a4+/+ or Tg(+);Slc26a4+/+ mice. No evidence of pendrin expression was found in the three sensory organs in Tg(+);Slc26a4+/− mice at P14 (Fig. S2A), P16 (Fig. 10A,C,D and Fig. S2C,G,K), P18 (S2E), and P35 (Fig. 10B). The absence of pendrin in Tg(+);Slc26a4+/− mice was observed with two different anti-pendrin antibodies (Pds1 and Pds2). In contrast, pendrin expression was found in transitional cells of the utricle, saccule and ampullae of controls that consisted of Tg(−);Slc26a4+/+ or Tg(+);Slc26a4+/+ mice. The expression patterns in Tg(−);Slc26a4+/+ and Tg(+);Slc26a4+/+ mice with both antibodies were similar to the pattern previously described in Slc26a4+/− mice [11,12].

Otoconia

Vestibular labyrinths were isolated by microdissection from Tg(−);Slc26a4+/−, Tg(+)−;Slc26a4+/− and Tg(+)−;Slc26a4+/− mice and the roof of the utricle was removed to permit an unobstructed view onto the utricular macula (Fig. 11A–C). Glittering otoconia were observed in Tg(+)−;Slc26a4+/− and Tg(+)−;Slc26a4+/− mice and giant otoconia in Tg(−);Slc26a4+/−. Otoconia were transferred into glass-bottom dishes and inspected by laser-scanning microscopy using a 405 nm laser. Giant otoconia from Tg(−);Slc26a4+/− mice were ~10-fold larger than normal otoconia (Fig. 11D). The shape of the giant otoconia resembled the shape previously observed in Slc26a4+/− mice [12]. Otoconia in Tg(+);Slc26a4+/− and Tg(+);Slc26a4+/− mice were similar consisting in both genotypes of larger (~20 μm, Figs. 11E,1 and F.1) and smaller (~10 μm; Figs. 11E,2 and F.2) otoconia, some of which revealed a concentric structure (Figs. 11E,3 and F.3). These data suggest that the introduction of the transgene rescued normal otoconia formation.

Balance

Balance tests were performed in Tg(+);Slc26a4+/− mice and Tg(+);Slc26a4+/− mice as well as in Tg(−);Slc26a4+/− and Tg(−);Slc26a4+/− mice (Fig. 12). Tg(−);Slc26a4+/− mice failed the test, which confirmed the vestibular phenotype previously described in Slc26a4+/− mice [14]. There was no apparent difference in the performance of Tg(+);Slc26a4+/−, Tg(+); Slc26a4+/− and Tg(−);Slc26a4+/− mice. These data demonstrate that the introduction of the transgene rescued normal gross motor vestibular function.

Discussion

In this study we generated a mouse that expresses human pendrin in the endolymphatic sac but lacks detectable pendrin protein expression in the cochlea or in the vestibular labyrinth. The most biologically interesting and clinically relevant observation of this study is that this mouse develops normal hearing and balance. Our findings support the hypothesis that pendrin expression in the endolymphatic sac is chiefly responsible for the development of normal endolymph volume, that lack of pendrin in the endolymphatic sac is mainly responsible for the development...
Figure 9. Hearing tests based on auditory brain stem responses. A–C) Examples of recordings of auditory brain stem responses to tone bursts of 16 kHz at amplitudes between 0 and 90 dB-SPL that were presented to Tg(+/);Slc26a4+/+ mice at P35 (A), Tg(−/);Slc26a4Δ/Δ mice at P35 (B), and Tg(+);Slc26a4Δ/+ mice at P35 (C). Thresholds at 15 dB-SPL are marked (thickened trace). D–L) Hearing thresholds based on auditory brain stem responses, were determined in Tg(+/);Slc26a4Δ/+ mice (D–F), Tg(+/);Slc26a4Δ/Δ mice (G–I), and Tg(+);Slc26a4Δ/+ mice (J–L; highlighted in red). Frequencies evaluated at 8 kHz (D, G, J), 16 kHz (E, H, K), and 32 kHz (F, I, L). The combined ranges of normal hearing in the three mouse strains (129S6, C57BL/6 and CBA) that contributed to the background of the mice are marked (grey rectangles). Numbers next to symbols in D–L represent the number of mice tested.

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Figure 10. Histology and pendrin expression in the vestibular labyrinth. Staining in all images consisted of immunocytochemistry of pendrin (Pds #1 antibody; red) and F-actin (green) and of nucleic acids (blue), the latter with the exception of images B and F. Images provide comparison of cryosections (A, C–D) and whole-mounted specimens (B) from Tg(+/)+Slc26a4+/− to cryosections (E, G–H) and whole-mounted specimens (F) from Tg(−)/Slc26a4−/− mice. Whole-mounted specimens were imaged by detecting fluorescence in 25 optical sections that were recorded in 1 µm intervals and projected into a single plane. The number of pairs of mice represented by these images are 2 for image A & E, 1 for B & F, 2 for C & G, and 2 for D & H with 3 sections being evaluated per animal. HC, vestibular hair cells; TC, transitional cells; M, melanocytes. Additional images using an alternative anti-pendrin antibody (Pds #2) and alternative positive controls (Tg(+/)+Slc26a4+/− mice) are provided in the Supplement (Fig. S2).

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of the membranous labyrinth enlargement in Slc26a4 mice, and that the complex inner ear pathology found in Slc26a4 mice is largely a consequence of the enlargement during embryonic development. This hypothesis was based on the studies in Slc26a4 mice that revealed that the enlargement is a key event on the path toward organ failure resulting in deafness and vestibular dysfunction [12,13,18] and on studies in Foxi1 mice that lack pendrin expression in the endolymphatic sac, develop an enlargement of the inner ear, but express pendrin in the cochlea and the vestibular labyrinth [19].

To test our hypothesis, we generated a transgenic mouse, Tg(B1-hPDS), which expresses human SLC26A4 (previously named PDS) controlled by the promoter of ATP1V1B1 [15]. The ability of the promoter of ATP1V1B1 to control gene expression had previously been evaluated in a transgenic mouse that expresses eGFP controlled by the promoter of ATP6V1B1 [16,20]. Expression of eGFP had been found in this mouse in intercalated cells of the renal collecting duct, and in narrow and clear cells of the epididymal epithelium of adult mice [16]. We found expression of eGFP in the embryonic kidney and in mitochon-

Figure 11. Otoconia. A–C) Microdissected maculae utriculi from a Tg(−);Slc26a4/ΔΔ, Tg(+/−);Slc26a4/ΔΔ and Tg(+/+);Slc26a4/ΔΔ mice. The maculae utriculi have been marked (red dashed line) and a single otoconium is indicated (red arrow head). D–F) Isolated otoconia from Tg(−);Slc26a4/ΔΔ, Tg(+/−);Slc26a4/ΔΔ and Tg(+/+);Slc26a4/ΔΔ mice. Note that images D, E.1, E.2, F.1 and F.2 are presented at the same scale. Mega-otoconia were found only in Tg(−);Slc26a4/ΔΔ mice. Tg(+/−);Slc26a4/ΔΔ mice contained a mixture of larger and smaller otoconia, some of which showed a concentric pattern (E.1, E.2 and E.3) that was similar to the mixture found in Tg(−);Slc26a4/ΔΔ mice (F.1, F.2 and F.3). doi:10.1371/journal.pgen.1003641.g011
dria-rich cells of the endolymphatic sac but not in the cochlea or the vestibular labyrinth (Fig. 3). Expression in mitochondria-rich cells of the endolymphatic sac was expected since these cells are members of the FORE family (forkhead related) of cells. FORE cells express FOXI1, which drives the expression of Atp6v1b1 and Slc26a4 [21,22,23]. Consistently, mitochondria-rich cells of the endolymphatic sac express the mRNAs Atp6v1b1 and Slc26a4 [24,25] and the corresponding proteins, the B1-subunit of the vH’ATPase and pendrin [11,23]. The onset of expression of Atp6v1b1 in the endolymphatic sac is at E11.5, which is similar to the onset of pendrin [13,26]. Thus, it was likely that the transgene Tg[B1-hPDS] would drive a timely expression of pendrin in mitochondria-rich cells of the endolymphatic sac.

Although FOXI1 drives the expression of Atp6v1b1 and Slc26a4 in FORE cells such as the mitochondria-rich cells of the endolymphatic sac, the expression of Atp6v1b1 and Slc26a4 is not limited to FORE cells. Indeed, Slc26a4 is expressed in the inner ear in spiral prominence and outer sulcus epithelial cells as well as in spindle-shaped cells of the cochlea and in transitional cells of the vestibular labyrinth, none of which are FORE cells [11,12]. Further, Atp6v1b1 expression has been found in the spiral limbs of the cochlea, which does not contain FORE cells [25,26]. The expression of Atp6v1b1 in the cochlea provided the possibility that the transgene would drive an ectopic expression of pendrin in the spiral limbus. Our studies of eGFP expression (Fig. 3) and of pendrin expression by Western blotting (Fig. 4) and immunocytochemistry of whole-mounted specimens and sections using two different anti-pendrin antibodies (Pds #1 and Pds #2, Fig. 7 and S1) revealed no detectable expression in the cochlea or vestibular labyrinth of Tg(+);Slc26a4Δ/Δ mice. The observed absence of pendrin expression in the vestibular labyrinth of Tg(+);Slc26a4Δ/Δ mice (Fig. 10 and S2) is consistent with the reported lack of Atp6v1b1 expression in the vestibular labyrinth based on detection by in situ hybridization [25,26] and by quantitative RT-PCR (Fig. 2B and F), which is a more sensitive technique. The observation that human SLC26A4 mRNA but no pendrin nor eGFP protein was detected in the cochlea or the vestibular labyrinth suggests the presence of strong translational regulation [27]. Taken together, our data demonstrate that we have generated a mouse that expresses pendrin in the endolymphatic sac but not in the cochlea or the vestibular labyrinth, although we cannot completely rule out that low levels of pendrin protein expression escaped our detection. Such low pendrin expression is unlikely the reason for the restored endolymphatic volume, since pendrin expression in the cochlea and vestibular labyrinth of Foxi1−/− mice, which lack pendrin expression in the endolymphatic sac, did not prevent endolymphatic enlargement [19] and since mice that express a mutant pendrin protein that supports anion exchange at a reduced rate are deaf, develop mega-otoconia and are balance impaired [28]. Moreover, hypomorphic mutant alleles of SLC26A4 show no difference in the resulting auditory phenotype from that of functional null alleles in patients with Pendred syndrome [29], indicating that small amounts of pendrin activity are insufficient to rescue hearing in humans.

Measurements of the endocochlear potential and pH revealed that the introduction of the transgene, which rescued normal endolymph volume, also rescued the loss of the normal endocochlear potential and the loss of the normal endolymphatic pH homeostasis (Fig. 8). It appears that the Cl−/HCO3− exchanger pendrin, which is normally expressed in the apical membranes of spiral prominence and outer sulcus epithelial cells, is not the sole mechanism responsible for the alkaline pH of endolymph in a normally developed cochlea. A similar conclusion can be drawn based on measurements in the doxycycline-inducible Slc26a4 mouse model where termination of pendrin expression at P6 led to the development of a nearly normal endocochlear potential and of a nearly normal alkaline pH [30]. We hypothesize that the epithelial barrier enclosing endolymph is permeable to H+, OH− and HCO3− and that the pH of endolymph follows the endocochlear potential.

Figure 12. Balance tests. Tests were based on the ability of Tg(+);Slc26a4Δ/Δ, Tg(+);Slc26a4Δ/Δ, Tg(−);Slc26a4Δ/Δ, Tg(−);Slc26a4Δ/Δ mice to balance on a rotating rod that was accelerated from 4 to 40 rpm over 60 s. Numbers near the symbols indicate the number of mice evaluated. doi:10.1371/journal.pgen.1003641.g012
Hearing and balance tests in Tg(+);Slc26a4+/A Δ mice revealed normal sensory function (Fig 9 and 12). The observation that hearing thresholds at 32 kHz had some variability in Tg(+);Slc26a4+/A Δ mice and that some Tg(+);Slc26a4+/A Δ mice developed progressive high-frequency hearing loss is most likely a function of the genetic background. Tg(+);Slc26a4+/A Δ mice were generated in a F2 generation from Slc26a4+/A Δ mice that were maintained isogenic in the 129S6 background and Tg(+);Slc26a4+/A Δ mice that were recently generated in a mixed background of C57BL/6 and CBA. DNA from the three background strains, 129S6, C57BL/6 and CBA, which differ in their hearing thresholds, are expected to comprise variable amounts of the genomes of individual mice. Hearing thresholds for 1 to 3 month-old 129S6, C57BL/6 and CBA mice range between 20–35 dB-SPL at 8 kHz, 10–28 dB-SPL at 16 kHz and 20–50 dB-SPL at 32 Hz [18,31,32,33,34,35]. In general, 129S6, C57BL/6 and CBA mice have similar thresholds at 8 kHz, whereas at 16 and 32 kHz CBA mice have lower thresholds than 129S6 and C57BL/6 mice. Thus, the greater variability in hearing thresholds that was observed at 32 kHz particularly in Tg(+);Slc26a4+/A Δ may be due to a variability in the mixture of these background strains.

Our observation that normal hearing developed in the absence of pendrin expression in the cochlea in combination with the published finding that normal hearing was maintained when pendrin expression was terminated after completed development [30], could suggest that pendrin in the cochlea has no physiologic significance beyond the developmental phase. However, it is also conceivable that pendrin-mediated HCO3− secretion provides a buffer that stabilizes the pH in the lateral wall tissues as well as in endolymph, and that this buffering is important during stress situations associated with normal life. Pendrin expression may indeed be important for the maintenance of hearing into advanced age.

In summary, we demonstrated that restoration of pendrin to the endolymphatic sac is sufficient to restore normal inner ear function. This implies that pendrin in the endolymphatic sac is more important for the development of normal hearing than pendrin expression in the cochlea and more important for the development of normal balance than pendrin expression in the vestibular labyrinth. This finding, in conjunction with our previous report that pendrin expression is required for embryonic development but not for the maintenance of hearing, opens the prospect that a spatially and temporally limited therapy will restore normal hearing in human patients carrying a variety of mutations of SLC26A4.

Methods

Ethics statement

All animal experiments and procedures at Kansas State University were performed according to protocols approved by the Animal Care and Use Committees at Kansas State University (IACUC#: 2961). All animal procedures at Sorbonne University Paris Cité were performed according to protocols approved by the ethics committee from University Pierre et Marie Curie, and were performed in accordance with the Guide for the Care and Use of Laboratory Animals (NIH publication No. 93-23, revised 1985).

Generation of Tg(B1-hPDS)+/+/Slc26a4+/+ transgenic mice

Human SLC26A4 cDNA was ligated into a pBluescript vector that contained 6.9 kb of the human ATP6V1B1 promoter [16,20]. An SV40 late region polyadenylation signal was cloned downstream of the SLC26A4 cDNA. The transgene Tg(B1-hPDS) included the 5′-flanking region of the ATP6V1B1 gene extending to but excluding the endogenous translational start codon, the human SLC26A4 cDNA, with its own translational start site, and the SV40 late region polyadenylation signal. The integrity of the transgene was confirmed by restriction digest and bidirectional sequencing of ligation sites. In preparation for injection, the transgene was linearized by SalI and NotI digestion, followed by gel purification using an electroelution method and then concentrated using EasiSpin columns (Whatman). The transgene was then further concentrated by ethanol precipitation and resuspended in low EDTA injection buffer (10 mM Tris with 0.1 mM EDTA). Tg(B1-hPDS) transgenic mice were created by the University of Utah transgenic mouse core facility using standard procedures [16,20]. Genotyping revealed that 63 pups were positive for transgene integration. One founder, which transmitted the transgene in a Mendelian fashion, was crossed with wild-type C57BL/6 × CBA F1 mice to establish a colony. Three Tg(B1-hPDS);Slc26a4+/+ transgenic mice were shipped to Kansas State University in Manhattan, Kansas, USA.

Generation of Tg(B1-hPDS)+/+/Slc26a4+/Δ transgenic mice

At Kansas State University, a colony of Tg(B1-hPDS)+/+, Slc26a4+/Δ mice was established. Colony management was supported by software (Litter tracker, written in Microsoft Visual Basic and Excel 2010 by P.W) Tg(B1-hPDS)+/+, Slc26a4+/Δ mice were crossed with Slc26a4+/Δ mice to generate Tg(B1-hPDS)+/+, Slc26a4+/Δ mice. Matings of Tg(B1-hPDS)+/+, Slc26a4+/Δ mice generated 28 Tg(B1-hPDS)+/+, Slc26a4+/Δ, 50 Tg(B1-hPDS)+/+, Slc26a4+/Δ and 39 Tg(B1-hPDS)+/+, Slc26a4+/Δ mice in a near Mendelian ratio of 1:2:1 with a 75% rate of transmission for the transgene (based on 169 pups).

Genotyping

Mice were genotyped for Slc26a4+/+ and Slc26a4−/− alleles by PCR using established primers [14] and for the transgene Tg(B1-hPDS) (Transnetyx, Cordova, TN). Primers for the transgene were designed to amplify a 345 bp PCR-product spanning the hPDS cDNA and the SV40 polyadenylation signal sequence (left primer: 5′-aggaagtcctttcaag-3′; right primer: 5′-cacaactagagtcatcg-3′) [15].

Isolation of embryonic tissues

Time-pregnant dams were deeply anesthetized with 4% tri-bromo-ethanol (0.014 ml/g body weight, i.p.) and embryos were harvested by laparotomy. Dams and embryos were sacrificed by decapitation. Gestational age was counted from the day when the vaginal plug was detected. This day was set to embryonic (E) day 0.5. Gestational age, however, was verified by evaluating gross morphological features including limbs, digits and the appearance of the pinna and auditory meatus [36,37].

Isolation of tissues from postnatal mice

The age of mice was counted from the day of birth, which was set to postnatal (P) day 0. Postnatal mice were deeply anesthetized with 4% tri-bromo-ethanol (0.014 ml/g body weight, i.p.) and sacrificed by decapitation or cardiac perfusion with fixative.

Quantitative RT-PCR

Quantitative RT-PCR was performed on total RNA [17]. Total RNA was isolated from tissues obtained by microdissection from Tg(−);Slc26a4+/− and Tg(+);Slc26a4+/−Δ mice and subjected to quantitative RT-PCR using gene-specific primers for 18S rRNA.
as well as for mRNA coding for the α-subunit of the mouse Na+/K+ ATPase Atp1a1, the β1-subunit of the mouse vH+ATPase Atp6v1b1, mouse pendrin Slc26a4 and human pendrin SLC26A4, which was introduced via the transgene.

Postnatal mice were genotyped by PCR prior to tissue collection. Embryonic Tg(−);Slc26a4TA/TA mice were generated by mating Tg(−);Slc26a4TA/TA dams and Tg(−);Slc26a4TA/TA sires, which yielded 100% of the desired genotype. Embryonic Tg(+/);Slc26a4TA/TA mice were generated by mating Tg(+/);Slc26a4TA/TA dams and sires, which yielded Tg(+/);Slc26a4TA/TA and Tg(−);Slc26a4TA/TA mice in a ratio of 3:1. Since embryonic mice could not be genotyped prior to tissue collection, the desired Tg(+/);Slc26a4TA/TA mice among Tg(−);Slc26a4TA/TA mice were initially identified by visual inspection of the size of the endolymphatic sac and the presence of ‘glittering’ otocoria. This phenotypic identification was subsequently confirmed by the presence of human pendrin SLC26A4 transgene by RT-PCR.

Tissues were obtained by microdissection. Endolymphatic sacs (8–10 endolymphatic sacs from 4–5 animals per sample) were obtained from mice at age E17.5. Cochlear ducts (4 cochlear ducts from 2 animals per sample and 2 cochlear ducts from 1 animal per sample) were obtained from mice at ages E17.5 and P2, respectively. Vestibular labyrinths (6 vestibular labyrinth samples from 2 animals per sample) were obtained from mice at age P8. Total RNA was isolated from microdissected tissues (RNaseasy micro kit, Qiagen, Valencia, CA, USA), treated with DNase (RNaseasy micro kit), combined with RNA storage solution (Applied Biosystems/Ambion, Austin, TX), adjusted to a concentration of 10 ng/μl, and stored at −80°C.

Quantity and quality of total RNA were evaluated by microfluidic electrophoresis (BioAnalyzer, Agilent, Santa Clara, CA), by microarray absorption photometry (Nanodrop, Wilmington, DE) and by quantitative RT-PCR of 18S rRNA. RNA was isolated from microdissected tissues (RNeasy micro kit, Qiagen, Valencia, CA, USA), treated with DNase (RNaseasy micro kit), combined with RNA storage solution (Applied Biosystems/Ambion, Austin, TX), adjusted to a concentration of 10 ng/μl, and stored at −80°C.

Efficiency

The number of template molecules (cDNA-template) was estimated to

\[
c_{\text{DNA template}} = \frac{6.02 \times 10^{23} \text{ molecules}}{\text{mol}} \times \frac{\text{ProductThreshold} \times \text{Efficiency}}{\text{ProductThreshold}} \times \text{Weight}_{\text{bp}} \times \frac{\text{mol}}{\text{bp}} \times \frac{\text{ProductSize} \times \text{bp}}{\text{mol}} \times \frac{1}{\text{Ct}}
\]

where 6.02×10^{23} molecules/mol represents Avogadro’s number, ProductThreshold is the weight of the PCR-product at threshold (0.49×10^{-3} g) that was obtained from calibration experiments, ProductSize is the size of the product in base pairs (bp), Weight_{bp} is average weight of one bp (660 g/mol), Efficiency is the PCR-efficiency obtained from the slope of the log-linear phase of the growth curve [38] and C is the cycle at which the fluorescense of the product molecules reaches a common threshold chosen in the middle of the log-linear part of the growth curve.

Paint-fill

Bisected heads of embryos age E15.5 were fixed overnight in Bodian’s fixatives, contained (vol/vol) 75% ethanol, 5% acetic acid, and 5% formalin in water. Heads were then dehydrated overnight in 100% ethanol and cleared in methyl-salicylate [39,40]. The membranous labyrinth was injected via the lateral wall of the basal turn of the cochlea and via the endolymphatic sac with diluted paint (Liquid Paper, Newell Rubbermaid, Atlanta, GA, 0.1–0.2% in methyl-salicylate) using a fine glass-electrode, a manipulator (NM-151 Narishige) and a micrometer-driven oil-filled microinjector (CellTram Vario, Eppendorf, Hamburg, Germany). For each genotype, at least three inner ears were injected.

Direct fluorescence of egFP

Whole mounts of fresh cochlear ducts, endolymphatic sacs and slices of kidney were prepared from E15.5 Tg(B1-egFP) mice and visualized with a fluorescence microscope (AxioScope, Carl Zeiss Witterscheid).

Western blotting

Crude membrane preparations of the inner ear. Soft tissues from 8 cochleae and 8 vestibular labyrinths devoid of endolymphatic sacs from adult mice were collected by microdissection in Ca2+-free solution and pooled and homogenized in 500 μl Tris-sucrose buffer. Ca2+-free solution contained (mM) 150 Na-gluconate, 1.6 K2HPO4, 0.4 KH2PO4, 4 Ca-gluconate, 2 MgSO4, and 5 glucose. pH 7.4. Tris-sucrose buffer contained (mM) 250 sucrose, 50 Tris-HCl, 5 EDTA, pH 7.4, and proteinase inhibitor cocktail (Cat# PI-87780 Fisher, Pittsburgh, PA). Homogenization involved a pestle/1.5 ml Eppendorf vial and a Potter-Evvelhjem tissue grinder (3 strokes at 1,250 rpm; Cat# K885512-0020, Fisher). Nuclei and debris were sedimented by centrifugation (6 min, 600 g, 4°C, Micromax RF, International equipment, Needham Heights, MA) and the supernatant contain-
ing membranes was collected. Pellets were washed 3 x with 100 µl Tris-sucrose buffer. Supernatants were pooled and membranes were pelleted by centrifugation (1 hr, 21,000 g, 4°C, Micromax RF). Membranes were suspended in 30 µl Tris-sucrose buffer and protein content was quantified (BCA assay, Cat# 23227, Fisher; Absorption photometer, Nanodrop).

**Crude membrane preparations of kidneys.** Kidneys (~1 g wet weight) were sliced, frozen in liquid N₂, pulverized, transferred into 3 ml Tris-sucrose buffer and homogenized. Homogenization involved pestles fitting 1.5 ml Eppendorf vials and the Potter-Elvehjem tissue grinder (3 strokes at 1,250 rpm). Nuclei and debris were sedimented by centrifugation (6 min, 600 g, 4°C, Micromax RF) and the supernatant containing membranes was collected. Pellets were washed 3 x with 2 ml Tris-sucrose buffer. Supernatants were pooled and membranes were pelleted by centrifugation (1 hr, 21,000 g, 4°C, Micromax RF). Membranes were suspended in 200 µl Tris-sucrose buffer and protein content was quantified (BCA assay).

**Gel-electrophoresis, blotting and protein detection.** Crude membranes were solubilized, denatured and reduced for 1 hr at 37°C in LDS sample buffer (Cat#: NP0007; Invitrogen) supplemented with 50 mM DTT. Proteins were resolved by gel-electrophoresis (4–12% Bis-Tris gels (Cat#: NP0326, Invitrogen) using a MOPS-SDS running buffer supplemented with an antioxidant (Cat#: NP0005, Invitrogen) and transferred onto PVDF membranes (BioRad, 3 hrs at 30 V). PVDF membranes were blocked for 1 hr at RT in TBST solution containing 3% BSA. TBST solution contained 20 mM Tris-base, 150 mM NaCl, 2.7 mM KCl, pH 7.4 with 0.15% Triton X 100) and 5% bovine serum albumin. Slides were incubated overnight at 4°C with primary antibody in PBS-TX with 2.5% BSA. Primary antibodies were two rabbit anti-pendrin antibodies (Pds #1; 1:200) and a polyclonal FITC-conjugated goat anti-eGFP antibody (Cat#: GTX26602, GeneTex, San Antonio, TX). Pds #1 was raised against the last 22 amino acids of the C-terminal of mouse pendrin. Human pendrin differs in 6 of these 22 amino acids [13,41]. Slides were washed with PBS-TX and, where appropriate, incubated for 1 hr at RT with secondary antibody (1:1,000, goat anti-rabbit Alexa 594, Invitrogen, Carlsbad, CA). After washing with PBS-TX, sections, where appropriate, were stained with phalloidin 488 (1:40; Invitrogen) to mark F-actin and DAPI (1:1,000; Invitrogen) to mark nuclei, washed again and cover-slipped with mounting medium (Vectashield HardSet Mounting Medium, Vector laboratories, Burlingame, CA).

**Whole-mounted specimens.** Tissues were obtained by microdissection from fixed temporal bones, washed in PBS-TX, and blocked with 3% bovine serum albumin in PBS-TX. Tissues were incubated overnight at 4°C with primary antibodies in PBS-TX with 2.5% BSA. Primary antibodies were a rabbit anti-pendrin antibody (Pds #1; 1:200) or a polyclonal FITC-conjugated goat anti-eGFP antibody (GeneTex). Slides were washed with PBS-TX and, where appropriate, incubated for 1 hr at RT with secondary antibody (1:1,000, goat anti-rabbit Alexa 594, Invitrogen). After washing with PBS-TX, tissues were stained with phalloidin 488 or phalloidin 594 (1:40; Invitrogen) and DAPI (1:1,000; Invitrogen), washed again and cover-slipped with mounting medium (Victashield HardSet Mounting Medium, Vector laboratories, Burlingame, CA).

**Confocal microscopy.** Immunocytochemistry of cryosections and whole-mounted specimens were viewed by confocal microscopy (LSM 510 Meta, Carl Zeiss, Göttingen, Germany). Laser scanning bright-field images were collected to aid orientation.

**Endocochlear potential and pH difference between endolymph and perilymph.**

Mice were anesthetized with 4% tri-bromo-ethanol for in situ measurements of the endocochlear potential and pH with double-barreled microelectrodes. Measurements were made in the basal turn of the cochlea by a round-window approach through the basilar membrane of the first turn of the cochlea [18,42]. The surgical cavity was covered with liquid Syglgard 184 (Dow Corning) to limit the loss of tissue CO₂ into ambient air.

Double-barreled glass microelectrodes were pulled (micropette puller PD-5; Narishige) from filament-containing glass tubing (1B100F-4; World Precision Instruments) and baked at 180°C for 2 h to ensure dryness. One barrel was silanized by a 30 s exposure to 0.008 ml dimethyldichlorosilane (40136; Fluka). After silanization, microelectrodes were baked again at 180°C for 2 h and tips were broken to a final O.D. of ~3 µm. The reference barrel was filled with 150 mM KCl and the ion-selective barrel was filled at the tip with liquid ion exchanger (Hydrogen ionophore II - Cocktail A, 95297; Fluka) and back-filled with buffer solution (500 mM KCl, 20 mM HEPES, pH 7.4).

Each barrel of the double-barreled microelectrode was connected via a Ag/AgCl₂ electrode to an electrometer (FD223, World Precision Instruments). A flowing KCl electrode (1 M KCl in 0.2% agar) was inserted under the skin of the animal to serve as ground electrode. Data were recorded in analog (BD12E Flattened recorder, Kipp & Zonen, Delft, The Netherlands) and digital form (DIGIDATA 1322A and AxoScope 10, Axon Instruments, Union City, CA). pH electrodes were calibrated in situ at 37°C using three
calibration solutions with different pH values. Calibration solutions contained (in mM): pH 6: 130 NaCl, 20 MES; pH 7: 130 NaCl, 20 HEPES; and pH 8: 130 NaCl, 20 HEPES. pH-sensitive electrodes had a slope of 56.9 ± 0.3 mV/pH unit (n = 11).

Hearing tests

Mice were deeply anesthetized with a mixture of dexmedetomidine and ketamine (0.375 mg/Kg body weight dexmedetomidine and 56 mg/kg body weight ketamine; i.p.) and placed on a thermal pad to maintain normal body temperature. The mastoid, vertex and ventral neck region of the animal were connected via sub-dermal platinum needle electrodes (TF-E2, Astro Med, Rhode Island, RI) and short (31 cm) leads to the main channel, reference channel and ground of the preamplifier, respectively. Auditory brainstem recordings were performed in a custom constructed, electrically shielded and sound-attenuated chamber (inner dimensions: 23 cm x 23 cm x 23 cm) using a digital data acquisition system (BioSig32 software, RA4LI Pre amplifier, RP2.1 Enhanced Real Time Processor, PA5 Programmable attenuator, ED1 Electrostatic Speaker Driver, Tucker-Davis Technologies, Alachua, FL). Tone burst stimuli were presented (21 per sec) via a free field electrostatic speaker (SigCal software, ES1 speaker, Tucker Davis). Acoustic stimuli were calibrated using a 1/4 inch condenser microphone (SigCal IRP4.2 software, Tucker Davis, PS9200 microphone, Acoustical Interface, Belmont, CA) placed at the location of the mouse head. Tone bursts (2 ms duration, 0.5 ms gate time; 8, 16, and 32 kHz) were presented with alternating phase (0 and 180°). Responses, recorded over 10 ms, were filtered (300 Hz high pass, 3000 Hz low pass and 60 Hz notch) and 1000 recordings were averaged. Tone burst stimuli were presented at intensities varying between 90 and 0 dB SPL in 5 dB intervals. Auditory thresholds were obtained by a visual comparison of wave forms. After the procedure, mice were rapidly recovered from anesthesia with atipamizole (1.875 mg/kg body weight; i.p.).

Balance tests

Balance testing consisted of determining the time that mice could balance on a rotating 1° rod with rotations ramping up from 4 to 40 rpm in 60 s (RotaRod, IITC Life Science, Woodland Hills, CA). Test chambers were cushioned with bubble-foil to provide a soft landing for mice falling off the rod.

Supporting Information

Figure S1 Histology and pendrin expression in the cochlea. Staining in all images consisted of immunocytochemistry of pendrin (Pds #1 antibody or Pds #2 antibody; red), F-actin (green) and nucleic acids (blue). Images provide comparison of pendrin expression in Tg(+);Slc26a4f/A mice (A, C, E) to positive controls consisting of Tg(+);Slc26a4f/A (B, D) or Tg(−);Slc26a4f/A mice (F). Note that both antibodies, Pds #1 and Pds #2, failed to detect pendrin expression in Tg(+);Slc26a4f/A mice and that the staining pattern for pendrin in positive controls was similar for both antibodies and for both positive controls. The number of pairs of mice represented by these images are 2 for images A & B, 1 for C & D, and 2 for E & F with 4 sections being evaluated per animal. K, Kölliker’s organ; OS, outer sulcus; LN, spiral lamina; IS, inner sulcus; OC, organ of Corti; SP, spiral prominence; SV, stria vascularis; RM, Reissner’s membrane. Compare these images to those in Fig. 7.

Figure S2 Histology and pendrin expression in the vestibular labyrinth. Staining in all images consisted of immunocytochemistry of pendrin (Pds #1 antibody or Pds #2 antibody; red) and F-actin (green) and of nucleic acids (blue). Images provide comparison of pendrin expression in Tg(+);Slc26a4f/A mice (A,C,E,G,I,K) to positive controls consisting of Tg(+);Slc26a4f/A (B, F, J) or Tg(−);Slc26a4f/A mice (D,H,L). Note that both antibodies, Pds #1 and Pds #2, failed to detect pendrin expression in Tg(+);Slc26a4f/A mice and that the staining pattern for pendrin in positive controls was similar for both antibodies and for both positive controls. The number of pairs of mice represented by these images are 2 for images A & B, 2 images C & D, 1 for images E & F, 2 for images G & H, 2 for images I & J, and 2 for images K & L. HC, vestibular hair cells; TC, transitional cells; M, melanocytes. Compare these images to those in Fig. 10.

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Author Contributions

Conceived and designed the experiments: PW RN NP RC DE TI AJG. Wrote the paper: XL PW. Mouse colony management and genotyping: TM.

References

1. Valvassori GE, Clemis JD (1978) The large vestibular aqueduct syndrome. Laryngoscope 88: 723–728.
2. Griffith AJ, Arts A, Downs C, Innis JW, Shepard NT, et al. (1996) Familial large vestibular aqueduct syndrome. Laryngoscope 106: 960–963.
3. Phelps PD, Coffey RA, Trembath RC, Luxon LM, Grossman AB, et al. (1998) Radiological malformations of the ear in Pendred syndrome. Clin Radiol 53: 269–273.
4. Griffith AJ, Wangemann P (2011) Hearing loss associated with enlargement of the vestibular aqueduct: Mechanistic insights from clinical phenotypes, genotypes, and mouse models. Hear Res 281: 11–17.
5. Ito T, Choi BY, King KA, Zalewski CK, Musket J, et al. (2011) SLC26A4 genotypes and phenotypes associated with enlargement of the vestibular aqueduct. Cell Physiol Biochem 28: 545–552.
6. Scott DA, Wang R, Kreman TM, Sheffield VC, Karninski LP (1999) The Pendred syndrome gene encodes a chloride-iodide transport protein. Nat Genet 21: 440–443.
7. Soleimani M, Greetley R, Petrovic S, Wang Z, Anklal H, et al. (2003) Pendrin: an apical CI−/OH−/HCO3− Exchanger in the Kidney Cortex. Am J Physiol Renal Physiol 280: F356–F364.
8. King KA, Choi BY, Zalewski C, Madeo AC, Manichaikul A, et al. (2010) SLC26A4 genotype, but not cochlear radiologic structure, is correlated with hearing loss in ears with an enlarged vestibular aqueduct. Laryngoscope 120: 384–389.
9. Takeda T, Taguchi D (2009) Aquaporins as potential drug targets for Meniere’s disease and its related diseases. Handb Exp Pharmacol 171–184.
10. Kim HM, Wangemann P (2010) Failure of fluid absorption in the endolymphatic sac initiates cochlear enlargement that leads to deafness in mice lacking pendrin expression. PLoS One 5: e14041.
12. Wangensteen P, Itza EM, Albrecht B, Wu T, Jabba SV, et al. (2004) Loss of KCNJ10 protein expression abolishes endocochlear potential and causes deafness in Pendred syndrome mouse model. BMC Medicine 2: 30
13. Kim HM, Wangensteen P (2011) Epithelial cell stretching and luminal acidification lead to a retarded development of stria vascularis and deafness in mice lacking pendrin. PLoS One 6: e17949.
14. Everett LA, Belyantseva IA, Noben-Trauth K, Cantos R, Chen A, et al. (2001) Expression of the forkhead transcription factor Foxi1 is a master regulator of vacuolar H+-ATPase proton pump subunits in the inner ear. Proc Natl Acad Sci U S A 98: 9727–9732.
15. Thibaut J, Picard N, Miller RL, Riemonsdy KA, Houiller P, et al. (2013) Mice with overexpression of the renal Cl-/HCO3- exchanger Pendrin (Slc26a4) develop chloride-sensitive hypertension. J Am Soc Nephrol. xxxx. In press.
16. Miller RL, Zhang F, Smith M, Beaulieu V, Paunescu TG, et al. (2005) V-ATPase B1-subunit promoter drives expression of CRE recombinase in intercalated cells of kidney, clear cells of epididymis and airway cells of lung in transgenic mice. Am J Physiol Cell Physiol 288: C1134–C1144.
17. Wangensteen P, Kim HM, Billings S, Nakaya K, Li X, et al. (2009) Developmental delays consistent with cochlear hypothyroidism contribute to failure to develop hearing in mice lacking Slc26a4/pendrin expression. Am J Physiol Renal Physiol 297: F1435–F1447.
18. Wangensteen P, Nakaya K, Wu T, Maganti R, Itza EM, et al. (2007) Loss of cochlear HCO3- secretion causes deafness via endolymphatic acidification and inhibition of Ca2+-reabsorption in a Pendred syndrome mouse model. Am J Physiol Renal Physiol 292: 1345–1355.
19. Hulander M, Kiernan AE, Blomqvist SR, Carlsson P, Samuelsson EJ, et al. (2007) Transcriptional control of SLC26A4 is involved in Pendred syndrome mouse model. Development 134: 2013–2025.
20. Miller RL, Lucero OM, Riemonsdy KA, Baumgartner BK, Brown D, et al. (2009) The V-ATPase B1-subunit promoter drives expression of CRE recombinase in intercalated cells of the kidney. Kidney Int 75: 435–439.
21. Blomqvist SR, Vidarsson H, Soder O, Enerback S (2006) Epifocal expression of the forkhead transcription factor Foxi1 is required for male fertility. EMBO J 25: 4131–4141.
22. Yang T, Vidarsson H, Rodrigo-Blaycayt S, Rosengren S, Enerback S, et al. (2007) Transcriptional control of SLC26A4 is involved in Pendred syndrome and nonsyndromic enlargement of vestibular aqueduct (DFN4). Am J Hum Genet 80: 1055–1063.
23. Vidarsson H, Westergren R, Heglid M, Blomqvist SR, Breton S, et al. (2009) The forkhead transcription factor Foxi1 is a master regulator of vacuolar H+-ATPase proton pump subunits in the inner ear, kidney and epididymis. PLoS One 4: e4741.
24. Everett LA, Morish H, Wu DK, Green ED (1999) Expression pattern of the mouse ortholog of the Pendred’s syndrome gene (Pds) suggests a key role for pendrin in the inner ear. Proc Natl Acad Sci U S A 96: 9727–9732.
25. Karet FE, Finberg KE, Nelson RD, Navir A, Mocan H, et al. (1999) Mutations in the gene encoding B1 subunit of H+-ATPase cause renal tubular acidosis with sensorineural deafness. Nat Genet 21: 84–90.
26. Dou H, Finberg K, Cartell DL, Laiou S, Choo D (2003) Mice lacking the B1 subunit of H+-ATPase have normal hearing. Hear Res 180: 76–84.
27. Hulander M, Enerback S (2000) Gene silencing by microRNAs: contributions of translational repression and mRNA decay. Nat Rev Genet 12: 99–110.
28. Dvor AA, Politi Y, Shahin H, Lenz DR, Dossena S, et al. (2010) Calcium oxalate stone formation in the inner ear-as a result of an Slc26a4 mutation. J Biol Chem 286: 21724–21735.
29. Choi BY, Stewart AK, Madeo AC, Pryor SP, Lenhard S, et al. (2009) Hyperfunctional SLC26A4 variants associated with nonsyndromic hearing loss and enlargement of the vestibular aqueduct: Genotype-phenotype correlation or coincidental polymorphisms? Hum Mutat 30: 599–608.
30. Choi BY, Kim HM, Ito T, Lee KY, Li X, et al. (2011) Mouse model of enlarged vestibular aqueduct defines temporal requirement of Slc26a4 expression for hearing acquisition. J Clin Invest 121: 4516–4525.
31. Zheng QY, Johnson K, Erway LC (1999) Assessment of hearing in 80 inbred strains of mice by ABR threshold analyses. Hear Res 130: 94–107.
32. Wilczyniller KK, Gagnon PM (2004) Cellular correlates of progressive hearing loss in 129/86/SvEv mice. J Comp Neurol 469: 377–390.
33. Ouagazzal AM, Reiss D, Romand R (2006) Effects of age-related hearing loss on stria vascularis and pre-epithelial inhibition in mice on pure and mixed C57BL and 129 genetic background. Behav Brain Res 172: 307–313.
34. Ison JR, Allen PD, O’Neil WE (2007) Age-related hearing loss in C57BL/6J mice has both frequency-specific and non-frequency-specific components that produce a hyperacusis-like exaggeration of the acoustic startle reflex. J Assoc Res Otorhinolaryngol 8: 539–550.
35. Frisina RD, Singh A, Bak M, Bozorg S, Seth R, et al. (2011) F1 (CBAxC57) mice show superior hearing in old age relative to their parental strains: hybrid vigor or a new animal model for “golden ears”? Neurobiol Aging 32: 1716–1724.
36. Rugh R (1968) The mouse: Its reproduction and development. Burgess Publishing Company. 430 p.
37. Theiler K (1972) The house mouse. Springer Verlag. 168 p.
38. Ramakers G, Ruijter JM, Deprez RH, Moorman AF (2003) Assumption-free analysis of quantitative real-time polymerase chain reaction (PCR) data. Neurosci Lett 339: 62–66.
39. Kiernan AE (2006) The pillafin method as a tool for analyzing the three-dimensional structure of the inner ear. Brain Res 297: 270–276.
40. Martin P, Swanson GJ (1993) Descriptive and experimental analysis of the mouse ortholog of the Pendred’s syndrome gene (Pds) suggests a key role for pendrin in the inner ear. Proc Natl Acad Sci U S A 96: 9727–9732.
41. Kim YH, Kwon TH, Frische S, Kim J, Tisher CC, et al. (2002) A new animal model for “golden ears”? Neurobiol Aging 32: 1716–1724.
42. Marcus DC, Rokugo M, Thalmann R (1985) Effects of barium and ion substitutions in artificial blood on endocochlear potential. Hear Res 17: 79–86.