The unique electrical properties in an extracellular fluid of the mammalian cochlea; their functional roles, homeostatic processes, and pathological significance

Fumiaki Nin 1 · Takamasa Yoshida 1,2,3 · Seishiro Sawamura 1 · Genki Ogata 1,2 · Takeru Ota 1 · Taiga Higuchi 1 · Shingo Murakami 4,5,6 · Katsumi Doi 7 · Yoshihisa Kurachi 4,5 · Hiroshi HIBINO 1,2,8

Received: 27 July 2016 / Accepted: 16 August 2016 / Published online: 27 August 2016 © The Author(s) 2016. This article is published with open access at Springerlink.com

Abstract The cochlea of the mammalian inner ear contains an endolymph that exhibits an endocochlear potential (EP) of +80 mV with a [K+] of 150 mM. This unusual extracellular solution is maintained by the cochlear lateral wall, a double-layered epithelial-like tissue. Acoustic stimuli allow endolym phatic K+ to enter sensory hair cells and excite them. The positive EP accelerates this K+ influx, thereby sensitizing hearing. K+ exits from hair cells and circulates back to the lateral wall, which unidirectionally transports K+ to the endolymph. In vivo electrophysiological assays demonstrated that the EP stems primarily from two K+ diffusion potentials yielded by [K+] gradients between intracellular and extracellular compartments in the lateral wall. Such gradients seem to be controlled by ion channels and transporters expressed in particular membrane domains of the two layers. Analyses of human deafness genes and genetically modified mice suggested the contribution of these channels and transporters to EP and hearing. A computational model, which reconstitutes unidirectional K+ transport by incorporating channels and transporters in the lateral wall and connects this transport to hair cell transcellular K+ fluxes, simulates the circulation current flowing between the endolymph and the perilymph. In this model, modulation of the circulation current profile accounts for the processes leading to EP loss under pathological conditions. This article not only summarizes the unique physiological and molecular mechanisms underlying homeostasis of the EP and their pathological relevance but also describes the interplay between EP and circulation current.

Keywords Cochlea · Hearing · Endocochlear potential · Ion concentrations · K+ transport · Electrophysiology · Computer simulation

Introduction

The auditory system continuously receives sounds of different intensities and frequencies from the ambient and extracts information necessary for the organism. The cochlea of the inner ear, the peripheral organ for hearing, harbors sensory hair cells that transduce mechanical stimuli to electrical signals. In humans, the audible frequencies range from 20 to 20,000 Hz, whereas the maximum hearing threshold is a trillionfold larger in acoustic power than the minimum threshold [44]. These striking profiles depend not only on the properties of hair cells but also on the ones of a unique extracellular solution that immerses them [14, 22, 36, 42, 109]. This solution is called “endolymph.”
The cochlea is filled with the following two different extracellular solutions: perilymph and endolymph. Textbooks for life science such as “Molecular Biology of the Cell” and “Principles of Neural Science” explain that, in general, extracellular solutions including blood plasma, interstitial fluid, and cerebrospinal fluid bear a $[K^+]$ of 5 mM and a $[Na^+]$ of 150 mM [3, 51]. This ionic composition is conserved in the perilymph. On the other hand, the cochlear endolymph contains 150 mM $[K^+]$ and 5 mM $[Na^+]$ (Fig. 1a) [91, 94]. Furthermore, this solution exhibits an endocochlear potential (EP) of +80 mV with respect to $K^+$ and 5 mM $Na^+$ (Fig. 1a) [91, 94]. Given that the stria harbors abundant vasculature, the EP greatly enhances these ionic flows by increasing the driving force and thereby crucially contributes to the achievement of superior profiles of hearing [14, 22, 42, 46]. Loss of EP causes deafness [42, 59]. After electrically exciting hair cells, $K^+$ is likely to exit from the basolateral membranes and reach the perilymph through $K^+$ channels (Fig. 1a) [20, 109].

The EP was measured for the first time in 1952 by Georg von Békésy, who won the Nobel Prize for the discovery of the physical mechanism of stimulation within the cochlea [5, 107]. Tasaki et al. (1959) found that the stria vascularis in the lateral cochlear wall is the source of the EP (Fig. 1a) [103]. Given that the stria harbors abundant vasculature, the maintenance of the EP was thought to need high energy. Indeed, anoxia causes a marked decrease in the EP [59, 63].

![Fig. 1](image-url)  
**Fig. 1** Electrochemical profile of the cochlea. a Structure of the cochlea. The composition of the human ear and a cross section of the cochlea are illustrated in the upper left and right panels, respectively. The lower panel depicts the tissue and cellular components of the cochlea. The upper panel shows the electrochemical properties of the endolymph and perilymph and the basolateral and apical membranes of the marginal cell layer, respectively. Shown in the lower panel is the EP endocochlear potential, OSC outer sulcus cells. Reproduced and modified from [74].

The composition of the endolymph and perilymph is conserved in the perilymph. Hair cells expose stereocilia extruded from their apical surface to the endolymph and bath their basolateral membranes in the perilymph. Acoustic stimuli open mechanoelectrical transduction (MET) channels at the top of the stereocilia, facilitating the entry of $K^+$ from the endolymph into hair cells. This $K^+$ influx depolarizes cells. Influx of $Ca^{2+}$, which is also contained at a concentration of approximately 20 μM in the endolymph, simultaneously occurs and tunes the hair cell functions involved in establishing the aforementioned dynamic ranges of audition [43]. The EP greatly enhances these ionic flows by increasing the driving force and thereby crucially contributes to the achievement of superior profiles of hearing [14, 22, 42, 46]. Loss of EP causes deafness [42, 59]. After electrically exciting hair cells, $K^+$ is likely to exit from the basolateral membranes and reach the perilymph through $K^+$ channels (Fig. 1a) [20, 109].

The EP was measured for the first time in 1952 by Georg von Békésy, who won the Nobel Prize for the discovery of the physical mechanism of stimulation within the cochlea [5, 107]. Tasaki et al. (1959) found that the stria vascularis in the lateral cochlear wall is the source of the EP (Fig. 1a) [103]. Given that the stria harbors abundant vasculature, the maintenance of the EP was thought to need high energy. Indeed, anoxia causes a marked decrease in the EP [59, 63].
that the extracellular space narrows down to 15 nm [41]. These membrane compartments are highly invaginated, so cell membranes and basolateral membranes of marginal cells. Characteristic is the complicated morphology of the intermediate from the two lymphs and the blood. Another important characteristic is the complex morphology of the intermediate cell membranes and basolateral membranes of marginal cells. These membrane compartments are highly invaginated, so that the extracellular space narrows down to 15 nm [41].

Classical pharmacological experiments using live animals characterized the channels and transporters expressed in the lateral wall and evaluated their possible physiological roles. Vascular and perilymphatic perfusion of ouabain, a blocker of Na⁺,K⁺-ATPases, or loop diuretics such as furosemide and bumetanide, antagonists for the Na⁺,K⁺,2Cl⁻-cotransporter (NKCC), dramatically suppressed the EP, suggesting that these K⁺-uptake transporters act in the stria vascularis and spiral ligament and critically contribute to the EP [38, 39, 62, 63, 66, 69, 90, 109]. Loss of EP was also observed upon systemic injection of Ba²⁺ that inhibits K⁺ channels, but it was not detected by perilymphatic application of this blocker [35, 70]. These results imply that the K⁺ channels expressed in the stria vascularis play key roles in the formation of the EP.

From the 1990s, advanced molecular biology, histochemical, and electrophysiological techniques have unveiled the molecular constituents and cellular and subcellular localizations of channels, transporters, and other apparatuses. It is likely that tight junctions in marginal cell layer contain claudin-1, -2, -3, -8, -9, -10, -12, and -14, and the ones in basal cells consist solely of claudin-11 [58]. Gap junctions included in the syncytial layer are constituted of connexin (Cx)-26, -29, -30, and -43 [55, 56, 67, 96, 114, 118]. Na⁺,K⁺-ATPases in the stria vascularis are complexes of α1 and β2 subunits and localize on the basolateral membranes of marginal cells, whereas those in the spiral ligament are composed of α1 and β1 subunits and are expressed on the membranes of fibrocytes [19, 73, 81]. The NKCC in the lateral wall is a type I cotransporter (NKCC1) and is present together with Na⁺,K⁺-ATPases in marginal cells and type II, IV, and V fibrocytes (Fig. 1b) [19, 73]. Furthermore, type I and III fibrocytes bear the K⁺,Cl⁻-cotransporter KCC3 [11]. We previously demonstrated that an inwardly rectifying K⁺ channel subunit KCNJ10 was expressed in the stria vascularis and responsible for Ba²⁺-sensitive K⁺ channels involved in the generation of EP [35]. Subsequently, another research group found that this channel subunit is localized on the membrane of intermediate cells (Fig. 1b) [4, 98]. In addition, in the marginal cells, the KCNQ1 and KCNE1 subunits were found to assemble and form functional IKs K⁺ channels (KCNQ1/KCNE1) on the apical membrane as well as Cl⁻ channels comprising the α subunit of CIC-Ka or CIC-Kb and the β subunit of barttin (CIC-K/barttin) on the basolateral membrane (Fig. 1b) [27, 77, 82, 88, 101]. The channels and transporters described above are thought to directly or indirectly contribute to unidirectional K⁺ transport across the lateral wall and to the EP formation. Moreover, patch-clamp assays of dissociated strial cells detected nonspecific cation (NSC) channels on the basolateral and apical membranes of marginal cells [95, 101, 102]. Apical channels comprise the TRPM4 subunit [83, 113].

On the lateral wall, other ion channels and transporters may be expressed and involved in the control of unidirectional K⁺ transport. Candidates have been obtained by our recent mass spectrometry analysis of membrane proteins extracted from rat stria vascularis [105]. Of 3236 proteins detected in the analysis, 16 ion channels and 62 transporters had not yet been identified in the stria. These include the K⁺ channels KCNJ13...
and KCNN4, as well as K⁺,Cl⁻-exchanger family proteins (KCCs). Although it must be carefully evaluated by histochemical approaches whether these K⁺ transport molecules are expressed in the stria vascularis or their detection by mass spectrometry results from the contamination from other tissues, their identification might reveal novel mechanisms of homeostasis of the EP.

**Pathological significance of the molecules expressed in the lateral wall**

Intensive research endeavors for the last 20 years have identified that genetic ablations and mutations of ion transport proteins expressed in the stria vascularis result in hearing impairment in mice or humans, as summarized in Table 1. For instance, KCNE1- or KCNQ1-null mice show severe collapse of the scala media and deafness [12, 106], implying that KCNQ1/KCNE1 K⁺ channels are crucial for driving K⁺ transport and maintaining the endolymph. These mice also show vestibular disorder. Loss-of-function mutations of the gene encoding both subunits cause Jarvell and Lange-Nielsen syndrome, accompanied by hearing disorders and arrhythmia. Similar phenotypes were observed in animals lacking the NKCC1-gene [24, 26, 31]. Type 4 Bartter’s syndrome, which is characterized by renal salt loss and hearing impairment, is induced by missense mutations in the amino-terminal region of the Cl⁻ channel subunit barttin [9, 27]. Knockout (KO) of this protein leads to loss of EP in mice [77]. KCNJ10-null mice display an ~0 mV EP with disruption of the structure of the scala media [71, 79]. Mutations in the gene of this channel subunit underlie the EAST/SeSAME syndrome, characterized by deafness, epilepsy, ataxia, mental retardation, tubulopathy, and electrolyte imbalance [10, 85]. Interest information has been further provided by analyses of the Na⁺,HCO₃⁻-exchanger SLC26A4, namely, pendrin, and the two-pore domain K⁺ (K₂P) channel subunit KCNK5. These two proteins are expressed in outer sulcus cells [13, 28, 78], which are also constituents of the lateral wall and localized beside the fibrocytes (Fig. 1a). Mutations in the SLC26A4-gene lead to Pendred syndrome, characterized by abnormal cochlear development, sensorineural hearing loss, and thyroid goiter [18, 49, 87]. Model mice of this disease display phenotypes of cochlear and vestibular dysfunctions with enlarged membranous labyrinths [28, 78]. KCNK5-KO mice exhibit loss of several cell types including outer sulcus epithelial cells and hair cells [13]. In these two mouse lines, the EP was markedly impaired. Therefore, outer sulcus cells may serve as a platform for ion transport involved in the maintenance of the endolymph.

| Molecules | Distribution | Human deafness types | Mouse phenotypes (other than deafness) | Reference |
|-----------|--------------|----------------------|----------------------------------------|-----------|
| KCNE1 (Isk, min K) | MC | Jarvell and Lange-Nielsen syndrome | Collapse of SM and VES, balance disorder | [106] |
| KCNQ1 (KvLQT1, Kv7.1) | MC | Jarvell and Lange-Nielsen syndrome | Collapse of SM and VES, balance disorder | [12] |
| KCNJ10 (Kir4.1) | IC | EAST syndrome, SeSAME syndrome | Collapse of SM, loss of EP (~0 mV) | [10, 71, 79, 85] |
| KCNK5 (K₃p5.1) | OSC | – | Loss of EP (~ +50 mV), degeneration of hair cells and spiral ganglions | [13] |
| BSND (barttin) | MC | Type 4 Bartter’s syndrome | Loss of EP (~ +17 mV) | [77] |
| NKCC1 (SLC12A1) | MC, FC | – | Collapse of SM and VES, balance disorder | [24, 26, 31] |
| Pendrin (SLC26A4) | SPEC, OSC | Pendred syndrome, DFNB4 | Loss of EP (~0 mV) | [2, 29, 30, 111] |
| SLC22A4 | StE | DFNB60 | – | [6] |
| KCC3 (SLC12A6) | FC | – | Degeneration of hair cells | [11] |
| GJA1 (Cx43) | FC | AR-nonsyndromic deafness | – | [68] |
| GJB1 (Cx32) | FC | X-linked Charcot-Marie-Tooth and deafness | – | [93] |
| GJB2 (Cx26) | FC | DFNB1, DFNA3A, hereditary palmoplantar keratoderma with deafness | Loss of EP (~ +50 mV), degeneration of hair cells | [16, 25, 54, 117, 119] |
| GJB3 (Cx31) | FC | DFNA2, AR-nonsyndromic deafness | – | [112] |
| GJB6 (Cx30) | FC | DFNA3B | Loss of EP (~ +3 mV), degeneration of hair cells | [34, 104] |
| Panx1 | FC, OSC | – | Loss of EP (~ +60 mV) | [15] |

**EP** endocochlear potential, **SM** scala media, **VES** vestibular endolymphatic space, **FCs** fibrocytes, **BCs** basal cells, **ICs** intermediate cells, **MCs** marginal cells, **OSCs** outer sulcus cells, **SPECs** spiral prominence epithelial cells, **StE** strial endothelium, **DFNA** the locus for autosomal dominant nonsyndromic deafness, **DFNB** the locus for autosomal recessive nonsyndromic deafness, **AR** autosomal recessive.
KO mice lacking any of the Cx26, 30, and 43 subunits show hearing loss, suggesting the importance of ion transport through the connexin network in the lateral wall [7, 8, 17, 16, 23, 32, 50, 52, 86, 104]. Finally, disruption of the claudin-11 gene results in deafness accompanied by a severe decrease in EP to \( \sim +30 \) mV [33, 57], and it highlights a crucial role for the barrier functions provided by tight junctions of basal cells in the stria vascularis.

**Mechanisms underlying the formation of the EP**

Along the identification of the molecular elements that contribute to the EP, the mechanisms underlying this process have been analyzed step by step. Pivotal observations were provided by in vivo electrophysiological experiments performed approximately 30 years ago. Salt et al. inserted a double-barreled K\(^+\)-selective microelectrode, which can simultaneously measure the potential and [K\(^+\)], from the perilymph toward the endolymph across the lateral wall in live guinea pigs [84]. In the stria vascularis, after detecting a cellular compartment exhibiting a high [K\(^+\)], they found a unique extracellular space called “intrastrial space (IS)” that showed a highly positive potential similar to the EP and a low [K\(^+\)] as observed in the perilymph. On the basis of these measurements, they assumed that (1) the cellular compartment close to the IS consisted of basal cells; (2) the IS potential (ISP) was primarily responsible for the EP and formed mainly by K\(^+\) diffusion potential, i.e., K\(^+\) equilibrium potential (\( E_K \)); and (3) this diffusion potential resulted from high K\(^+\) permeability on the apical surfaces of basal cells. The electrochemical properties of the IS were also confirmed by another report [45]. Later on, patch-clamp and immunohistochemical experiments demonstrated that K\(^+\) permeability dominated the membranes of intermediate cells, as it emerged from KCNJ10 and other K\(^+\) channels (Fig. 1b) [4, 100]. According to the aforementioned histological studies [41], the IS corresponds to the 15-nm extracellular space primarily surrounded by invaginated membranes of intermediate and marginal cells (Fig. 1b) [84].

By incorporating these observations, the “two-cell model” and the “five-compartment model” have been proposed to explain the generation of the EP [84, 108]. In this model, the EP represents a potential difference across the lateral wall composed of two layers,

\[
EP = v_{SB} - v_{SB} = v_{MB} - v_{MA},
\]

where \( v_{SB} \) and \( v_{SA} \) are the membrane potentials across the basolateral and apical surfaces of the syncytial layer, respectively, and \( v_{MB} \) and \( v_{MA} \) are the ones across the basolateral and apical surfaces of the marginal cell layer, respectively (Fig. 1b, c). Each membrane potential was relative to the neighboring extracellular solution, which was defined as 0 mV. Equation 1 indicates that the EP depends on the four membrane potentials provided by the two layers of the lateral wall. Given that K\(^+\) conductance governs the intermediate cell membranes that correspond to the apical surface of the syncytial layer [100], Eq. 1 can be described by the equation

\[
EP = \frac{RT}{F} \ln \left( \frac{[K^+]_{IS}}{[K^+]_{SY}} \right) + v_{MB} - v_{MA}. \tag{2}
\]

In this scenario, the ISP is equivalent to the transepithelial voltage across the syncytial layer,

\[
ISP = v_{SB} - v_{SA} = v_{SB} - v_{MB} - \ln \left( \frac{[K^+]_{IS}}{[K^+]_{SY}} \right). \tag{3}
\]

According to patch-clamp analyses and in vivo experiments [72, 84, 99], \( v_{SB}, v_{MB}, \) and \( v_{MA} \) were estimated to be small of \(<10 \) mV [100]. Therefore, it was assumed that the values of \( v_{SA}, ISP, \) and EP are approximately equal to each other (Eqs. 2 and 3). This relationship can be described as follows [100]:

\[
EP = \frac{RT}{F} \ln \left( \frac{[K^+]_{IS}}{[K^+]_{SY}} \right) \approx ISP, \tag{4}
\]

where \([K^+]_{IS}\) and \([K^+]_{SY}\) are the \([K^+]\) in the IS and inside the syncytial layer, respectively. We assayed the lateral wall of live guinea pigs with double-barreled K\(^+\)-selective microelectrodes and confirmed the aforementioned potential and \([K^+]\) in the IS, as shown in Fig. 1c [1, 74, 116]. Our measurements also determined the properties of other compartments. First, the potential inside the syncytial layer, which is equivalent to \( v_{SB} \) in Eqs. 1–3, is moderately positive (\( \sim +7 \) mV), compared to the perilymph. Second, there is only a slight transepithelial voltage across the marginal cell layer (i.e., \( v_{MB} - v_{MA} \); see Eqs. 1 and 2), and thereby, the ISP is similar to the EP. These observations reinforce the idea that the ISP represents the K\(^+\) diffusion potential and dominates the EP, as described by Eq. 5.

Under physiological conditions, \([K^+]_{SY}\) greatly exceeds \([K^+]_{IS}\) (Fig. 1c), providing a highly positive potential (\( \sim +80 \) mV) for the ISP and EP (Eq. 4). On the other hand, when Na\(^+\), K\(^+\)-ATPases or NKCC in the stria vascularis were impaired by anoxia, which would mimic ischemia, or by vascular perfusion of blockers, the EP dramatically reduced to \(30 \) to \(40 \) mV [59]. Under these conditions, it is plausible that \([K^+]_{SY}\) is smaller than \([K^+]_{IS}\) (Eq. 4) or that \( v_{SB}, v_{MB}, \) or \( v_{MA} \) is significantly modulated (Eq. 2). However, these alterations were not considered in the two-cell model or five-compartment model. These issues were addressed by our fine in vivo measurements [74]. In the example shown in Fig. 2, a K\(^+\)-selective microelectrode was held in the IS, while the other electrode was being inserted in the endolymph to continuously
monitor the EP. Imposition of anoxia on the animal caused an increase in $[K^+]_{IS}$ and a decrease in ISP. Importantly, the ISP calculated from Eq. 3 with the measured $[K^+]_{IS}$, $[K^+]_{SY}$, and $v_{SB}$ (constant) was in agreement with the ISP directly recorded with the $K^+$-selective microelectrode. This finding supports that the ISP stems primarily from the $K^+$ diffusion potential. On the other hand, the simultaneously measured EP was reduced more strongly than the ISP and reached a negative value. This indicates that anoxia can enlarge the transepithelial voltage across the marginal cell layer ($v_{MB} - v_{MA}$) and that this property is involved in the reduction of the EP. To clarify the underlying mechanisms, we examined marginal cells and found that a large $K^+$ diffusion potential was induced on the apical surface of the marginal cell layer and it formed the difference between the EP and ISP (Figs. 3A, Ba). Under physiological conditions, the marginal cell $K^+$ diffusion potential, which is likely to emerge from KCNQ1/KCNE1 channels (Fig. 1b), is relatively small, because of the similarity between $[K^+]$ in the marginal cell layer ($[K^+]_{MC}$) and $[K^+]$ in the endolymph ($[K^+]_{EL}$) (Fig. 3A). Application of anoxia markedly decreased $[K^+]_{MC}$ with a minimal effect on $[K^+]_{EL}$, increasing the diffusion potential (Fig. 3Ba) [74]. Little change was detected in the membrane potential across the basolateral surface of the marginal cell layer ($v_{MB}$), in spite of the elevation of $[K^+]_{IS}$ and the reduction of $[K^+]_{MC}$. This experimental observation is reasonable, given that patch-clamp analysis of isolated marginal cells detected abundant $Cl^-$ conductance but little $K^+$ conductance on their basolateral membranes [88]. Similar results were obtained when ouabain or the loop diuretic bumetanide was vascularly applied (Fig. 3Ba) [74]. In summary, the EP primarily depends on two $K^+$ diffusion potentials on the apical surfaces of the syncytial and marginal cell layers,

$$EP \approx v_{SB} + \frac{RT}{F} \ln \left( \frac{[K^+]_{IS}}{[K^+]_{SY}} \right) + v_{MB} - \frac{RT}{F} \ln \left( \frac{[K^+]_{MC}}{[K^+]_{EL}} \right) \quad (5)$$

**Integration of circulation current into processes for formation of the EP by theoretical approaches**

Under physiological conditions, $[K^+]_{IS}$ is lower than $[K^+]_{MC}$ (Figs. 1c and 3A). However, blocking $Na^+,K^+$-ATPases or NKCC1 in marginal cells, $[K^+]_{IS}$ and $[K^+]_{MC}$ gradually change, and finally, the former exceeds the latter (Fig. 3Ba) [37, 74]. These $[K^+]$ dynamics, which underlie the reduction of ISP and EP, may derive from the modulation of currents through channels and transporters in the lateral wall. Nevertheless, such currents are neither measurable directly by in vivo electrophysiological approaches nor described in previous models and equations. Therefore, clarification of this issue requires theoretical approaches.

Each of the marginal cell and syncytial layers expresses particular $K^+$ channels or $K^+$-uptake transporters on different surfaces (Fig. 1b). Assembly of these two layers expresses abundant $Cl^-$ channels but little $K^+$ conductance on their basolateral membranes [88]. Similar results were obtained when ouabain or the loop diuretic bumetanide was vascularly applied (Fig. 3Ba) [74]. In summary, the EP primarily depends on two $K^+$ diffusion potentials on the apical surfaces of the syncytial and marginal cell layers,
endolymph across the lateral wall. MET channels in hair cells are slightly open even in their resting state [80], so that in vivo K⁺ constantly flows through cells. Taking these considerations together, it is plausible that, even without acoustic stimulation, K⁺ is transported from the perilymph to the endolymph and then returns to the endolymph through the three layers in the cochlea, i.e., the syncytial, marginal cell, and hair cell layers (Fig. 1a). This “circulation current” composed of K⁺ [75], also referred to as “K⁺ recycling” [55, 56, 109, 110], likely corresponds to an idling current in the electrical circuit. The existence of a circulation current was first proposed in Davis’ battery theory [21], and it was validated by in vivo recording of the voltage gradient representing ion flows with a microelectrode in the perilymph [120]. The circulation current has been suggested to be involved in the maintenance of high potential and [K⁺] in the endolymph [20, 55, 56, 109, 110].

The conditions of cells controlled by circulation current are different from the ones of cells isolated from the organism. When isolated cells are bathed in artificial extracellular solution filling an experimental chamber, their resting membrane potential (RMP) can be predicted with the Nernst or Goldman-Hodgkin-Katz (GHK) equations. In this “equilibrium state,” the total current across the cell membrane is zero. This is the principle of the two-cell model or five-compartment model (Eqs. 1–4) [20, 84, 100] as well as the one of our earlier model (Eq. 5) [37, 74]. On the other hand, in vivo, the unidirectional circulation current is likely to constantly flow across the three layers even under physiological conditions. In this case, membrane potentials as well as ion concentrations in every cell are in the “steady state” and therefore constant. Based on this idea, we developed a computational model, the “Ninh-Hibino-Kurachi (NHK) model,” to quantitatively describe the dynamics of the electrochemical properties of the cochlea (Fig. 4A) (for detailed equations, see [75]). In this model, the profile of each membrane is characterized by equivalent circuits of Hodgkin-Huxley-type equations that represent the function of channels and transporters. The circuits in six membrane domains were then connected in series to reconstitute the circulation current. In other words, the circulation current corresponds to the sum of the currents through all the channels and transporters in each membrane domain. This arrangement allows the circulation current to control membrane potentials as well as ion concentrations in the extracellular and intracellular compartments. In particular, the latter parameter is described by the following equation:

$$\frac{d[X]}{dt} = \frac{I_{X, in} - I_{X, out}}{V \times F},$$  (6)

where $[X]$ is the concentration of ion X, $I_{X, in}$ is the inward current of X, $I_{X, out}$ is the outward current of X, V is the volume of the extracellular or intracellular compartment, and F is the Faraday constant. Each of $I_{X, in}$ and $I_{X, out}$ is the sum of the current passing through an X-permeable channel and the X-flow through the transporter carrying X on the membrane. Note that $I_{X, in}$ and $I_{X, out}$ are components of the circulation current. Finally, the current passing through MET channels always equals the circulation current in the route across the hair cell layer. Thus, the amplitude of the circulation current depends on the potential difference across the apical surface of hair cells, i.e., the difference between the EP and the potential.
inside hair cells with respect to the perilymph. To summarize, under physiological conditions, the amplitude of the circulation current is constant and flows through every membrane of the three layers; \( I_{X_{in}} \) is same as \( I_{X_{out}} \) for every compartment (Eq. 6). This situation provides constant values for all of the ionic concentrations (\( \frac{d[X]}{dt} = 0 \)), all of the membrane potentials, and the EP. However, when functions of channels or transporters are perturbed, the circulation current changes, modulating ionic concentrations and membrane potentials.

By using the NHK model, we simulated the electrochemical profile in the lateral wall and endolymph under physiological conditions and during anoxia that mimics ischemic conditions. Our in vivo experiments strongly suggest that imposition of anoxia on animals inhibited not only Na\(^+\),K\(^+\)-ATPases but also NKCC on the basolateral surface of the marginal cell layer [75]. The measured behaviors for EP, ISP, \([K^+]_{IS}\), and \([K^+]_{MC}\) were highly reproducible in silico, when the activities of both of these \( K^+ \)-uptake transporters were reduced to 5% of the control (Fig. 4B). Simulated changes of


### Fig. 4

Computer simulation of electrochemical dynamics in the cochlea.

**A** Key elements in the Nin-Hibino-Kurachi (NHK) model. The profile of each membrane is described by electrical circuits that represent the function of channels and transporters. The circuits in six membrane domains deriving from the syncytial, marginal cell, and hair cell layers were connected in series to allow the flow of circulation current, which is carried by K⁺ under physiological conditions. NKCC1 Na⁺,K⁺-cotransporter type 1, CIC CIC-K/barttin type Cl⁻ channel, NSC nonselective cation channel, TJ tight junction, MET mechanoelectrical transduction, baso basolateral, api apical. **B** Reproduction of electrochemical properties in the cochlea under physiological and anoxic conditions. Change of endocochlear potential (EP; 

It is surprising that in vivo and under physiological conditions, the basolateral surface of the syncytial layer exhibits a significantly positive potential (≈ +7 mV) with respect to the perilymph (v_SB ≈ +7 mV; Figs. 1c and 3A) [116], although in general, eukaryotic cells show a negative membrane potential at their resting state [40, 48, 76]. The positive RMP of the syncytial layer is critical for harvesting the K⁺ diffusion potential on the apical surface to set the ISP and EP to a highly positive value (Eqs. 3 and 5), which is essential for hearing. As already mentioned, the syncytial basolateral surface is mainly provided by fibrocytes in the spiral ligament. In spite of the physiological importance of fibrocytes, the machineries underlying the establishment of this unique RMP have not yet been characterized. A major reason is likely to be the technical difficulty of patch-clamp assays conducted on fibrocytes tightly embedded in the extracellular matrix [53]. Our in vivo experiments with double-barreled K⁺-selective microelectrodes demonstrated that v_SB is remarkably hyperpolarized by perilymphatic perfusion of a solution with low [Na⁺] (Fig. 3Bd), but it can only moderately be changed by a solution containing either high [K⁺] or low [Cl⁻] [115]. These observations imply that the fibrocyte membrane is more permeable to Na⁺ than to K⁺ and Cl⁻ (Fig. 3A), and this unusual current properties in the lateral wall are schematically illustrated in Fig. 4C [75]. Under physiological conditions, the circulation current is set to be purely made up of K⁺ at any membrane domain (Fig. 4Ca). During anoxia, marginal cell transporters, which are blocked as mentioned above, can barely transport K⁺. Instead, the circulation current can pass as flows of Cl⁻ and Na⁺ through the CIC-K/barttin and NSC channels, respectively; both of which are coexpressed with the two K⁺-uptake transporters. This indicates that the carrier of the circulation current can switch from K⁺ to other ions on the basolateral surface of the marginal cell layer (Fig. 4Cb). Consequently, K⁺ influx exceeds K⁺ efflux in the IS and vice versa in the marginal cell layer, causing an increase in [K⁺]IS and a decrease in [K⁺]MC (Fig. 4Bb and Bc). In the late phase of anoxia, [K⁺]IS significantly exceeds [K⁺]MC (Fig. 3Bb). These changes gradually affect membrane properties and therefore reduce the ISP and EP (Fig. 4Ba). This EP impairment further results in the reduction of the amplitude of the circulation current.

### Profile of the basolateral surface of the syncytial layer

The EP is determined by the potentials across the four membrane domains of the lateral wall, i.e., v_SB, v_SA, v_MB, and v_MA (see Eqs. 1 and 5 and Fig. 3A, Ba). As described above, the marginal cell apical and basolateral surfaces as well as the syncytial layer apical surface, which corresponds to the intermediate cell membrane, have been so far considerably characterized by multiple approaches. However, little is known about the functional elements of the syncytial layer basolateral surface, formed primarily by fibrocytes of the spiral ligament [108]. Earlier studies demonstrated that perfusion of either ouabain or loop diuretics to the perilymph, in which fibrocytes are immersed, dramatically reduces the EP [69, 90]. Therefore, among several proteins expressed in fibrocytes, Na⁺,K⁺-ATPases and NKCC seem to regulate electrochemical properties and K⁺ transport in the lateral wall (Fig. 1b). This idea was tested in our in vivo experiments with K⁺-selective double-barreled microelectrodes. Perilymphatic application of ouabain markedly decreased [K⁺]SY with a minimum change of [K⁺]IS [1] (Fig. 3Bb). As a consequence, the ISP and the EP declined (Eqs. 3 and 5). On the other hand, whereas bumetanide was perfused to the perilymph, [K⁺]SY barely changed [116] (Fig. 3Bc). These results strongly suggest that, in the syncytial layer, Na⁺,K⁺-ATPases, but not NKCC1, functionally transport K⁺ across the basolateral surface and maintain a high [K⁺]SY, essential for the large K⁺ diffusion potential on the apical surface. Further analyses revealed that during perilymphatic perfusion of bumetanide, [K⁺]IS increases and [K⁺]MC decreases, as observed during vascular perfusion [116] (Fig. 3Bc). Therefore, this compound is likely to reach the NKCC1 on the basolateral surface of the marginal cell layer and impair the ISP and EP in processes similar to anoxia (see Eq. 5 and Fig. 3Ba).
profile is critically involved in establishment of the positive RMP under physiological conditions. During perfusion of the low-Na+ solution, the ISP and EP were reduced to an extent similar to the alternation of $v_{SB}$ (Fig. 3Bd), suggesting that this perturbation minimally affects the other membrane potentials in the lateral wall.

Conclusions and perspectives

As described above, various experimental techniques have demonstrated the molecular and physiological architectures that drive unidirectional K+ transport across the lateral wall. This ionic flow is likely to be involved in maintaining the electrochemical properties of the syncytial and marginal cell layers and control the two K+ diffusion potentials that form the EP. Correlation of the circulation current, which underlies K+ transport across the lateral wall, with the EP has also been demonstrated by theoretical approaches. In spite of these achievements, several issues remain to be clarified. First, it will be necessary to identify the proteins responsible for Na+ permeability, which plays a key role in maintaining the positive RMP in fibrocytes. A second interesting objective would be to identify the triggers that activate the fibrocyte NKCC1, a transporter that is likely to be silent under physiological conditions. The third open question is the process by which KCC3 in fibrocytes as well as SLC26A4 and KCNK5 in outer sulcus cells regulate the EP. Fourth, some of the channels and transporters in the lateral wall are expected to be functionally affected by Ca2+ or H+, but the relevance of these ions to the EP is still elusive. This issue may require detailed information on several Ca2+ or H+ transporting molecules detected by our mass spectrometry experiments conducted on strial proteins [105]. All of these analyses will enable us to improve the NHK model and thereby more precisely describe how the EP and circulation current interplay with each other. A combination of improved computational models and conventional experimental approaches will help us to theoretically understand the reactions of the EP and the properties of the lateral wall in response to different acoustic stimuli, as well as to identify the pathophysiological mechanisms underlying hearing impairment observed in various human diseases and animal models. Further research will contribute to advance the auditory science and medical therapies for hearing disorders.

Acknowledgments This work was partially supported by the following research grants and funds: Grant-in-Aid for Scientific Research C 15K10770 (to KD); Grants-in-Aid for Young Scientists A 15H05683 (to FN); Grants-in-Aid for Scientific Research on Innovative Areas “HD Physiology” 25136704 (to FN) and 22136002 (to YK) from the Ministry of Education, Culture, Sports, Science, and Technology of Japan; AMED-CREST, AMED (to HH); a Grant for Promotion of Niigata University Research Projects 24A006 (to HH); grants from The Salt Science Research Foundation Nos. 1318 and 1549 (to FN); The Takeda Science Foundation (to FN); The Uehara Memorial Foundation (to FN); and Astellas Foundation for Research on Metabolic Disorders (to FN).

Compliance with ethical standards

Ethical standards The experiments comply with the current laws of Japan.

Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

References

1. Adachi N, Yoshida T, Nin F, et al. (2013) The mechanism underlying maintenance of the endocochlear potential by the K+ transport system in fibrocytes of the inner ear. J Physiol 591:4459–4472. doi:10.1113/jphysiol.2013.258046
2. Albert S, Blons H, Jonard L, et al. (2006) SLC26A4 gene is frequently involved in nonsyndromic hearing impairment with enlarged vestibular aqueduct in Caucasian populations. Eur J Hum Genet 14:773–779. doi:10.1038/sj.ejhg.5201611
3. Alberts B, Johnson A, Lewis J, Raff M, Roberts K, Walter P (2014) Molecular biology of the cell, 5th edn. New York press, New York
4. Ando M, Takeuchi S (1999) Immunological identification of an inward rectifier K+ channel (Kir4.1) in the intermediate cell (melanocyte) of the cochlear stria vascularis of gerbils and rats. Cell Tissue Res 298:179–183
5. Bekesy G (1952) DC resting potentials inside the cochlear partition. J Acoust Soc Am 24:72–76
6. Ben Said M, Grati M, Ishimoto T, et al. (2016) A mutation in SLC22A4 encoding an organic cation transporter expressed in the cochlea strial endothelium causes human recessive nonsyndromic hearing loss DFNB60. Hum Genet 135:513–524. doi:10.1007/s00439-016-1657-7
7. Berger AC, Kelly JJ, Lajoie P, et al. (2014) Mutations in Cx30 that are linked to skin disease and nonsyndromic hearing loss exhibit several distinct cellular pathologies. J Cell Sci 127:1751–1764. doi:10.1242/jcs.138230
8. Bicego M, Beltramello M, Melchionda S, et al. (2006) Pathogenetic role of the deafness-related M34 T mutation of Cx26. Hum Mol Genet 15:2569–2587. doi:10.1093/hmg/ddl184
9. Birkenhager R, Otto E, Schurmann MJ, et al. (2001) Mutation of BSND causes Bartter syndrome with sensorineural deafness and kidney failure. Nat Genet 29:310–314. doi:10.1038/ng752
10. Bockenhauer D, Feather S, Stanesco HC, et al. (2009) Epilepsy, ataxia, sensorineural deafness, tubulopathy, and KCNJ10 mutations. N Engl J Med 350:1960–1970. doi:10.1056/NEJMoA0810276
11. Boettger T, Rust MB, Maier H, et al. (2003) Loss of K-Cl co-transporter KCC3 causes deafness, neurodegeneration and reduced seizure threshold. EMBO J 22:5422–5434. doi:10.1093/emboj/cdg519
12. Casimiro MC, Knollmann BC, Ebert SN, et al. (2001) Targeted disruption of the Kcnq1 gene produces a mouse model of Jervell and Lange-Nielsen syndrome. Proc Natl Acad Sci U S A 98:2526–2531. doi:10.1073/pnas.041398998
intercellular gap junctional coupling. J Assoc Res Otolaryngol 13: 473–484. doi:10.1007/s10162-012-0322-7

54. Kelsell DP, Dunlop J, Stevens HP, et al. (1997) Connexin 26 mutations in hereditary non-syndromic sensorineural deafness. Nature 387:80–83. doi:10.1038/387080a0

55. Kikuchi T, Adams JC, Miyabe Y, et al. (2000) Potassium ion recycling pathway via gap junction systems in the mammalian cochlea and its interruption in hereditary nonsyndromic deafness. Med Electron Microsc 33:51–56. doi:10.1021/000000009

56. Kikuchi T, Kimura RS, Paul DL, et al. (1995) Gap junctions in the rat cochlea: immunohistochemical and ultrastructural analysis. Anat Embryol (Berl) 191:101–118

57. Kitajiri SI, Furuse M, Morita K, et al. (2004) Expression patterns of claudins, tight junction adhesion molecules, in the inner ear. Hear Res 187:25–34

59. Konishi T, Butler RA, Fernandez C (1961) Effect of anoxia on cochlear potentials. J Acoust Soc Am 33:349–356

60. Konishi T, Hamrick PE, Walsh PJ (1978) Ion transport in guinea pig cochlea. I. Potassium and sodium transport. Acta Otolaryngol 86:22–34

61. Konishi T, Kelsey E (1973) Effect of potassium deficiency on cochlear potentials and cation contents of the endolymph. Acta Otolaryngol 76:410–418

62. Konishi T, Mendelsohn M (1970) Effect of ouabain on cochlear potentials and endolymph composition in guinea pigs. Acta Otolaryngol 69:192–199

63. Kuipers W, Bonting SL (1970) The cochlear potentials. I. The effect of ouabain on the cochlear potentials of the guinea pig. Pflugers Arch 320:348–358

64. Kuipers W, Bonting SL (1970) The cochlear potentials. II. The nature of the cochlear endolymphatic resting potential. Pflugers Arch 320:359–372

65. Kumar NM, Ghalila NB (1996) The gap junction communication channel. Cell 84:381–388

66. Kusakari J, Rokugo M, Kambayashi J, et al. (1983) Generation of endocochlear potential depends on two K + diffusion potentials and an electrical barrier in the stria vascularis of the inner ear. Proc Natl Acad Sci U S A 105: 1751–1756. doi:10.1073/pnas.0711461015

67. Nin F, Hibino H, Doi K, et al. (2008) The endocochlear potential depends on two K + diffusion potentials and an electrical barrier in the stria vascularis of the inner ear. Proc Natl Acad Sci U S A 105: 1751–1756. doi:10.1073/pnas.0711461015

68. Liu XZ, Xia XJ, Adams J, et al. (2001) Mutations in GJA1

69. Marcus DC, Demott JE, Kobayashi T, et al. (2001) Connexin 26 mouse mutant represents a model for the human hereditary keratitis-ichthyosis-deafness syndrome. Hum Mol Genet 10:2945–2951

70. Marcus DC, Rokugo M, Thalmann R (1985) Effects of barium and beta subunit isoforms of the Na,K-ATPase in the rat cochlea. Mol Cell Neurosci 2:179–187

71. Marcus DC, Wu T, Wangemann P, et al. (2002) KCNJ10 (Kir4.1) and beta subunit isoforms of the Na,K-ATPase in the rat cochlea. J Comp Neurol 438:216–230

72. Melichar I, Syka J (1987) Electrophysiological measurements of the stria vascularis potentials in vivo. Hear Res 25:165–172

73. Nakazawa K, Spiker SS, Schulte BA (1995) Ultrastructural localization of Na,K-ATPase in the gerbil cochlea. J Histochem Cytochem 43:981–991

74. Nin F, Hibino H, Doi K, et al. (2008) The endocochlear potential depends on two K + diffusion potentials and an electrical barrier in the stria vascularis of the inner ear. Proc Natl Acad Sci U S A 105: 1751–1756. doi:10.1073/pnas.0711461015

75. Nin F, Hibino H, Murakami S, et al. (2012) Computational model of a circulation current that controls electrochemical properties in the mammalian cochlea. Proc Natl Acad Sci U S A 109:9191–9196. doi:10.1073/pnas.120067109

76. Noble D (1984) The surprising heart: a review of recent progress in cardiac electrophysiology. J Physiol 353:1–50

77. Rickheit G, Maier H, Strenzeke N, et al. (2008) Endocochlear potential depends on Cl− channels: mechanism underlying deafness in Barter syndrome IV. EMBO J 27:2907–2917. doi:10.1038/emboj.2008.203

78. Royaux IE, Belyantseva IA, Wu T, et al. (2003) Localization and functional studies of pendrin in the mouse inner ear provide insight into the etiology of deafness in Pendred syndrome. J Assoc Res Otolaryngol 4:394–404

79. Rozengurt N, Lopez I, Chiu CS, et al. (2003) Time course of inner ear degeneration and degeneration in mice lacking the Kir4.1 potassium channel subunit. Hear Res 177:71–80

80. Russell JJ, Cady AR, Richardson GP (1986) The responses of inner and outer hair cells in the basal turn of the guinea-pig cochlea and in the mouse cochlea grown in vitro. Hear Res 22:199–216

81. Ryan AF, Watts AG (1991) Expression of mRNA encoding alpha and beta subunit isoforms of the Na,K-ATPase in the rat cochlea. Mol Cell Neurosci 2:179–187

82. Sakagami M, Fukazawa K, Matsunaga T, et al. (1991) Cellular localization of rat Isk protein in the stria vascularis by immunohistochemical observation. Hear Res 56:168–172

83. Sakuraba M, Murata J, Teruyama R, et al. (2014) Spatiotemporal expression of TRPM4 in the mouse cochlea. J Neurosci Res 92: 1409–1418. doi:10.1002/jnr.23410

84. Salt AN, Melichar I, Thalmann R (1987) Mechanisms of endocochlear potential generation by stria vascularis. Laryngoscope 97:984–991

85. Salt AN, Melichar I, Thalmann R (1987) Mechanisms of endocochlear potential generation by stria vascularis. Laryngoscope 97:984–991

86. Schutz M, Auth T, Gehrt A, et al. (2011) The connexin26 S17F channel in excised membrane patches of strial marginal cells. Laryngoscope 121:1720–1724

87. Sheffield VC, Kraiem Z, Beck JC, et al. (1996) Pendred syndrome maps to chromosome 7q21-34 and is caused by an intrinsic defect in thyroid iodine organification. Nat Genet 12:424–428. doi:10.1038/ng0496-424

88. Shen Z, Marcus DC (1998) Divalent cations inhibit Isk/KvLQT1 channels in excised membrane patches of stria marginal cells. Hear Res 123:157–167

89. Shindo M, Miyamoto M, Abe N, et al. (1992) Dependence of endocochlear potential on basolateral Na+ and Cl− concentration: a study using vascular and perilymph perfusion. Jpn J Physiol 42: 617–630

90. Shugyo A, Mori N, Matsunaga T (1990) A comparison of the concentration-dependent deafness in X-linked Charcot-Marie-tooth disease. Hum Mol Genet 10:2945–2951

91. Smith CA, Lowry OH, ML W (1954) The electrolytes of the labyrinthine fluids. Laryngoscope 64:141–153. doi:10.1289 /0005537-19540300-00001

92. Sohl G, Maxeiner S, Willecke K (2005) Expression and functions of neuronal gap junctions. Nat Rev Neurosci 6:191–200. doi:10.1038/nrn1627

93. Stojkovic T, Latour P, Vandenbergh A, et al. (1999) Sensorineural deafness in X-linked Charcot-Marie-tooth disease with connexin 32 mutation (R142Q). Neurology 52:1010–1014
94. Suga F, Nakashima T, JB S Jr (1970) Sodium and potassium ions in endolymph. In vivo measurements with glass microelectrodes. Arch Otalaryngol 91:37–43

95. Sunose H, Ikeda K, Saito Y, et al. (1993) Nonselective cation and Cl⁻ channels in luminal membrane of the marginal cell. Am J Physiol 265:C72–C78

96. Suzuki T, Takamatsu T, Oyamada M (2003) Expression of gap junction protein connexin43 in the adult rat cochlea: comparison with connexin26. J Histochem Cytochem 51:903–912

97. Takeuchi S, Ando M (1998) Dye-coupling of melanocytes with endothelial cells and pericytes in the cochlea of gerbils. Cell Tissue Res 293:271–275

98. Takeuchi S, Ando M (1998) Inwardly rectifying K⁺ currents in marginal cells of stria vascularis. Hear Res 61:86

99. Takeuchi S, Ando M, Irimajiri A (1997) Changes in the volume of intermediate cells in the cochlea of gerbils: a possible contribution to the endocochlear potential. Neurosci Lett 258:9–12

100. Takeuchi S, Ando M, Kakigi A (2000) Mechanism generating endocochlear potential: role played by intermediate cells in stria vascularis. Biophys J 79:2572–2582. doi:10.1016/S0006-3495(00)76497-6

101. Takeuchi S, Ando M, Kozakura K, et al. (1995) Ion channels in basolateral membrane of marginal cells dissociated from gerbil stria vascularis. Hear Res 83:89–100

102. Takeuchi S, Marcus DC, Wangemann P (1992) Ca²⁺-activated nonselective cation, maxi K⁺ and Cl⁻ channels in apical membrane of marginal cells of stria vascularis. Hear Res 61:86–96

103. Tasaki I, Spyropoulos CS (1959) Stria vascularis as source of nonselective cation, maxi K⁺ and Cl⁻ channels in apical membrane of marginal cells. Proc Natl Acad Sci USA 45:1369–1374

104. Teubner B, Michel V, Pesch J, et al. (2003) Connexin30 (Gjb6)-deficiency causes severe hearing impairment and lack of endocochlear potential. Hum Mol Genet 12:13–21

105. Uetsuka S, Ogata G, Nagamori S, et al. (2015) Molecular architecture of the stria vascularis membrane transport system, which is essential for physiological functions of the mammalian cochlea. Eur J Neurosci 42:1984–2002. doi:10.1111/ejn.12973

106. Vetter DE, Mann JR, Wangemann P, et al. (1996) Inner ear defects induced by null mutation of the Isk gene. Neuron 17:1251–1264

107. Von Bekesy G (1952) Resting potentials inside the cochlear partition of the guinea pig. Nature 169:241–242

108. Wangemann P (1995) Comparison of ion transport mechanisms between vestibular dark cells and strial marginal cells. Hear Res 90:149–157

109. Wangemann P (2002) K⁺ cycling and the endocochlear potential. Hear Res 165:1–9

110. Wangemann P (2006) Supporting sensory transduction: cochlear fluid homeostasis and the endocochlear potential. J Physiol 576:11–21. doi:10.1113/jphysiol.2006.112888

111. Wangemann P, Itza EM, Albrecht B, et al. (2004) Loss of KCNJ10 protein expression abolishes endocochlear potential and causes deafness in Pendred syndrome mouse model. BMC Med 2:30. doi:10.1186/1741-7015-2-30

112. Xia JH, Liu CY, Tang BS, et al. (1998) Mutations in the gene encoding gap junction protein beta-3 associated with autosomal dominant hearing impairment. Nat Genet 20:370–373. doi:10.1038/3945

113. Yamaguchi S, Tanimoto A, Otsuguro K, et al. (2014) Negatively charged amino acids near and in transient receptor potential (TRP) domain of TRPM4 channel are one determinant of its Ca²⁺ sensitivity. J Biol Chem 289:35265–35282. doi:10.1074/jbc.M114.606087

114. Yang JJ, Liao PJ, CC S, et al. (2005) Expression patterns of connexin29 (GJE1) in mouse and rat cochlea. Biochem Biophys Res Commun 338:723–728. doi:10.1016/j.bbrc.2005.09.193

115. Yoshida T, Nin F, Ogata G, et al. (2015) NKCCs in the fibrocytes of the spiral ligament are silent on the unidirectional K⁺ transport that controls the electrochemical properties in the mammalian cochlea. Pflugers Arch 467:1577–1589. doi:10.1007/s00424-014-1597-9

116. Yoshida T, Nin F, Murakami S, et al. (2016) The unique ion permeability profile of cochlear fibrocytes and its contribution to establishing their positive resting membrane potential. Pflugers Arch. 468:1609–1619. doi:10.1007/s00424-016-1853-2

117. Zelante L, Gasparini P, Estivill X, et al. (1997) Connexin26 mutation associated with Pendred syndrome mouse model. BMC Med 2:30. doi:10.1186/1741-7015-2-30

118. Zhao HB, Kikuchi T, Ngezahayo A, et al. (2006) Gap junctions and cochlear homeostasis. J Membr Biol 209:177–186. doi:10.1007/s00232-005-0832-x

119. Zhu Y, Chen J, Liang C, et al. (2015) Connexin26 (GJB2) deficiency reduces active cochlear amplification leading to late-onset hearing loss. Neuroscience 284:719–729. doi:10.1016/j.neuroscience.2014.10.061

120. Zidanic M, Brownell WE (1990) Fine structure of the intracochlear potential field. I. The silent current. Biophys J 57:1253–1268. doi:10.1016/S0006-3495(90)82644-8