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Published in:
Frontiers in Neurology

DOI:
10.3389/fneur.2011.00048

2011

Citation for published version (APA):

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Identification of new signaling components in the sensory epithelium of human saccule

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Objective: To locate components and target proteins of relevance for the cAMP and cGMP signaling networks including cAMP and cGMP phosphodiesterases (PDEs), salt-inducible kinases (SIKs), subunits of Na+, K+-ATPases, and aquaporins (AQPs) in the human saccule.

Methods: The human saccule was dissected out during the removal of vestibular schwannoma via the translabyrinthine approach and immediately fixed. Immunohistochemistry was performed using PDE, SIK, Na+, K+-ATPase, and AQP antibodies. Results: PDEs selective for cAMP (PDE4A, PDE4D, and PDE8A) and cGMP (PDE9A) as well as a dual specificity PDE (PDE10A) were detected in the sensory epithelium of the saccule. Furthermore, AQP2, 4, and 9, SIK1 and the α-1 subunit of the Na+, K+-ATPase were detected. Conclusion: cAMP and cGMP are important regulators of ion and water homeostasis in the inner ear. The identification of PDEs and SIK1 in the vestibular system offers new treatment targets for endolympathic hydrops. Exactly how the PDEs are connected to SIK1 and the SIK1 substrate Na+, K+-ATPase and to AQPs 2, 4, 9 remains to be elucidated. The dissection of the signaling networks utilizing these components and evaluating their roles will add new basic knowledge regarding inner ear physiology.

Keywords: saccule, immunohistochemistry, cAMP, cGMP, cyclic nucleotide phosphodiesterase, salt-inducible kinase, Na, K-ATPase, aquaporin

INTRODUCTION

The membranous labyrinth of the inner ear is a sensory system for sound, motion, and gravity, consisting of the cochlea, vestibular system, and the endolymphatic sac. The lumen of the membranous labyrinth is filled with endolymph, a K+-rich, positively polarized fluid, whereas the surrounding spaces are filled with perilymph, with a composition similar to regular extracellular fluid (Andrews, 2004; Thalmann et al., 2006; Lang et al., 2007). Dysregulation of ion and water homeostasis in the inner ear is believed to result in endolympathic hydrops, a condition associated with vertigo and hearing loss (Semaan et al., 2005). Several studies indicate an important role for the cAMP second messenger system in the regulation of ion and water homeostasis in the inner ear. For example, cAMP has been shown to regulate the secretion of K+ into the endolymph (Wangemann, 2002; Salt and Plontke, 2010) and it has been suggested that water homeostasis in the inner ear is regulated in part via the vasopressin–cAMP–aquaporin (AQP)2 water channel system (Takeda and Taguchi, 2009) in the same fashion as in the kidney (Lang et al., 2007; Noda et al., 2010). When it comes to the cGMP signaling system and the regulation of ion and water homeostasis in the inner ear, less is known. However, roles for the nitric oxide–cGMP and the atrial natriuretic peptide (ANP)–cGMP systems have been suggested (Fessenden and Schacht, 1998; Semaan et al., 2005; Borghì et al., 2006). ANP has hypotensive and hypovolemic effects which are mediated via increases in intracellular cGMP levels (Ahlawalia et al., 2004). Hypotension has been suggested to play a role in inner ear disorders (Pirodda et al., 1997, 2001) and ANP receptors are expressed in the inner ear (Long et al., 2010).

By hydrolyzing cAMP and cGMP, cyclic nucleotide phosphodiesterases (PDEs) regulate a wide variety of biological responses mediated by these second messenger molecules. Mammalian PDEs can be sorted into 11 functionally distinct, highly regulated, and structurally related families (Manganiello et al., 2006; Conti and Beavo, 2007). These PDE families differ in their primary sequences, substrate affinities, and catalytic properties, sensitivity to effectors and inhibitors, responses to regulatory molecules, and cellular functions. Some PDE families are specific for cAMP hydrolysis (PDEs 4, 7, 8), others are cGMP-specific (PDEs 5, 6, 9), and some hydrolyze both cGMP and cAMP (PDEs 1, 2, 3, 10, 11). Most cells contain representatives of more than one PDE gene family, but in different amounts, proportions, and subcellular locations. By virtue of their distinct intrinsic characteristics and their intracellular targeting to different subcellular locations, different PDEs integrate multiple cellular inputs and modulate the amplitude, duration, termination, and specificity of cyclic nucleotide signals and actions (Manganiello et al., 2006; Conti and Beavo, 2007; Houslay, 2010).

Very little is known about PDEs and how they relate to other signaling networks and targets in the inner ear. In this study we...
Table 1 | Antibodies: source, type, immunogen, positive/negative controls.

<table>
<thead>
<tr>
<th>Antibody/source/cat #/antibody type</th>
<th>Immunogen</th>
<th>Positive/negative control</th>
</tr>
</thead>
<tbody>
<tr>
<td>AQP2 (H-40)/Santa Cruz/sc 28629/rabbit polyclonal</td>
<td>Amino acid 232–291 within the C-terminal domain of human AQP2</td>
<td>Ref1/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>AQP4 (H-80)/Santa Cruz/sc 20812/rabbit polyclonal</td>
<td>Amino acid 244–323 within the C-terminal domain of human AQP4</td>
<td>Refs2,3/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>AQP9/Abcam/ab84828/rabbit polyclonal</td>
<td>A 15 amino acid synthetic peptide from the N-terminal of human AQP9</td>
<td>Stains AQP9 using immunohistochemistry on human liver sections (Abcam)/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>PDE4A, PDE4D, PDE8A, PDE9A, PDE10A/Scottish Biomedical/rabbit polyclonal</td>
<td>Designed from the C-terminal region from respective human enzyme</td>
<td>Refs4,5 (PDEs 8A, 9A, 10A) and Omar et al. (unpublished) (PDE4A and 4D)/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>N^+/K^+ ATPase α-1/Millipore/05-369/mouse monoclonal</td>
<td>Purified N^+,-K^+ ATPase isolated from membrane fractions of rat kidney outer medulla</td>
<td>Refs6,7/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>SIK1 (Y-20)/Santa Cruz/sc83754/rabbit polyclonal</td>
<td>A peptide within an internal region of human SIK1</td>
<td>Refs8,9/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>Caveolin-1 (N-20)/Santa Cruz/sc-894/rabbit polyclonal</td>
<td>A peptide at the N-terminal of human caveolin-1</td>
<td>Refs10/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>Laminin γ1/Gift Dr. Sorokin/mouse monoclonal</td>
<td>Rat glomerular basement membrane membrane</td>
<td>Ref11/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>Pan-laminin 1/485/Gift Dr. Sorokin/rabbit polyclonal</td>
<td>Mouse laminin α1, β1, γ1</td>
<td>Ref12/omission of the antibody (no reaction)</td>
</tr>
<tr>
<td>Gial fibrillary acidic protein/Sigma-Aldrich/G3893/mouse monoclonal</td>
<td>Purified GFAP from pig spinal cord</td>
<td>Ref13/omission of the antibody (no reaction)</td>
</tr>
</tbody>
</table>

1Maekawa et al., 2010 (human endolymphatic sac); 2Ishiyama et al., 2009 (human utricle); 3Kakigi et al., 2009 (human endolymphatic sac); 4Heimann et al., 2010 (human pancreatic islets); 5Omar et al., accepted for publication (human adipose tissue); 6Kobayashi et al., 2009 (rat erythrocytes); 7Yang et al., 2011 (mouse coelomic, stria vascularis capillaries); 8Noin et al., 2009 (mouse liver); 9Nilsson et al., 2006 (rat adipocytes); 10Hoé et al., 2009 (rat astrocytes); 11Sanes et al., 1990 (muscle fiber, nerve trunk, kidney, blood vessels from rat, rabbit, guinea pig and human); 12Ishiyama et al., 2010 (human utricle); 13Agrawal et al., 2006 (mouse brain).
Invitrogen) was added for 10 min to visualize all cell nuclei. At the end of the incubation, the sections were washed in PBS and mounted with Vectashield (Vector Laboratories). Sections were observed using a camera-equipped fluorescence microscope. Images were obtained using a Zeiss Axioshot 2 microscope and a Hamamatsu C4742-95 camera with OpenLab 5 software (Improvision) for image processing. The quality of the microdissected saccule was determined by staining with hematoxylin and eosin. For all staining, several sections from three saccules were analyzed.

RESULTS
To map the expression pattern of PDEs and other signaling components and targets of relevance for the cAMP and cGMP signaling networks and for the regulation of water and ion homeostasis, an immunohistochemistry approach was used.

Figure 1A shows hematoxylin and eosin staining of a representative section of human saccule. The sensory epithelium with nuclei at several layers, non-sensory epithelium, otoconia, the endolymph compartment, as well as the stroma, with cells, can be seen. Glial fibrillary acidic protein (GFAP) antibodies were used to stain supporting cells in the sensory epithelium (Figures 1B–D; Lopez et al., 2007). Figure 1B is an overview showing the sensory epithelium and the non-sensory epithelium. Figures 1C,D show supporting cells as well as hair cells.

As shown by representative photographs (Figures 2A–J), the sensory epithelium of the saccule expresses PDEs specific for cAMP (PDE4A, PDE4D, and PDE8A), for cGMP (PDE9A), and a dual specificity enzyme (PDE10A). The immunoreactivity to PDE10A was mainly found in the apical part of the epithelium (Figures 2I,J). As is shown in Figures 2A,B, no significant staining in the non-sensory epithelium was obtained for PDE4A. This was also the case for the other PDEs investigated (data not shown).

The water channel AQP2 is known to be regulated by vasopressin in a CAMP-dependent manner (Takeda and Taguchi, 2009) as has been investigated in detail in kidney cells (Lang et al., 2007). Immunoreactivity to AQP2 is expressed in the sensory epithelium of the human saccule as well as in stromal cells (Figures 3A,C) but not in the non-sensory epithelium (data not shown). The expression of AQP4 and AQP9, two other AQPs known to be regulated in cAMP- and cGMP-dependent manners, were then investigated. Immunoreactivity to AQP4 was detected in the basolateral part of the sensory epithelium (Figures 3D–F) as well as in non-sensory epithelium (data not shown). Basolateral location of AQP4 was confirmed by co-expression with laminin 1, a basal membrane protein (Figures 3G,H). In previous studies AQP4 has been shown to co-stain with flotillin-1 in brain astrocytes, indicating raft location of AQP4 (Noel et al., 2009). We found no staining for flotillin-1 (unpublished observation) in human saccule sensory epithelium, however, immunoreactivity of another raft marker, caveolin-1, was detected in the sensory epithelium (Figure 3B) but not in the non-sensory epithelium (data not shown). Immunoreactivity to AQP9 was detected partially in the basolateral part of the epithelium as indicated by co-stained with laminin 1. AQP9 immunoreactivity was not seen in the non-sensory epithelium (Figures 3I,J).

The expression of SIK1, another CAMP-regulated target of relevance for ion and water homeostasis, was investigated. Immunoreactivity to SIK1 was detected in the sensory epithelium (Figures 4A,B). SIK1 has been shown to regulate the activity of Na+, K+-ATPase in kidney cells (Iaitovich and Bertorello, 2010; Taub et al., 2010), and AQP4 has been shown to associate with a Na+, K+-ATPase (Illarionova et al., 2010). The α-1 subunit of the Na+, K+-ATPase was expressed in the human sacculus epithelium and immunostaining was partially localized to the basolateral part of the cells as shown by co-staining with laminin 1 (Figures 5A–C).

Figure 6 shows a schematic view of the possible interrelation of signaling components and targets studied in this work. Included in the figure is also CAMP-dependent protein kinase (PKA) and cGMP-dependent protein kinase (PKG), the major mediators of CAMP and cGMP effects, respectively. PDEs with the function to specifically catalyze the hydrolysis of cAMP and cGMP in response to signaling inputs, such as vasopressin, catecholamines, atrio natriuretic peptide, and nitric oxide may be important in the regulation of water and ion homeostasis in the inner ear by having impact on SIK1 and the SIK1 substrate Na+, K+-ATPase as well as on a number of AQPs. The presence of these signaling networks in the inner ear needs to be verified and studied at the functional level.

DISCUSSION
cAMP and cGMP have important roles in the regulation of water and ion homeostasis in the inner ear (Fessenden and Schacht, 1998; Wangemann, 2002; Semaan et al., 2005; Salt and Plontke, 2010). Regulation of the cyclic nucleotides occurs at the level of synthesis...
by adenylate and guanylate cyclases as well as at the level of degradation by PDEs. PDEs have been studied in many different cell types and these have been shown to express different members of the 11 PDE families (Manganiello et al., 2006; Conti and Beavo, 2007). Here we show for the first time that PDE proteins are expressed in the inner ear. More specifically, we demonstrate the expression of PDEs selective for cAMP (PDE4A, PDE4D, PDE8A) and for cGMP (PDE9A) as well as of a PDE hydrolyzing both

**FIGURE 2 | Immunostaining of phosphodiesterases (red) in human saccules.** Merged photos of PDE4A, laminin (green), and DAPI (blue (A)), merged photos of PDE4A and DAPI (B), PDE4D (C), merged photos of PDE4D and DAPI (D), PDE8A (E), merged photos of PDE8A and DAPI (F), PDE9A (G), merged photos of PDE9A and DAPI (H), PDE10A (I), merged photos of PDE10A, laminin (green), and DAPI (J). Sensory epithelium (bar with two arrowheads), transition to non-sensory epithelium (vertical bar with one arrow head), lumen (lu), stroma (st). Bars 30 μm (A,B), 20 μm (C–J).

**FIGURE 3 | Immunostaining of aquaporins (red).** AQP2 (A), merged photos of AQP2 and DAPI (blue (C)), Caveolin-1 (red (B)), AQP4 (D,E), merged photos of AQP4 and DAPI (F), merged photos of AQP4 and laminin (green (G)), laminin (green (H)), merged photos of AQP9, laminin and DAPI (I), and merged photos of AQP9 and laminin (J). Lumen (lu), bm [basal membrane (G–J)], stroma (st), sensory epithelium (bar with two arrow heads), stromal cell (A,C, bar with one arrow head), transition to non-sensory epithelium (I,J, bar with one arrow head). Bars 15 μm (A,C), 12.5 μm (B), 25 μm (D), 15 μm (E–H), 20 μm (I,J).
cAMP and cGMP (PDE10A) in the sensory epithelium of human saccule. These enzymes could have a number of physiological roles when it comes to the regulation of cAMP- and cGMP-mediated processes in the saccule related to water and ion homeostasis. For example, PDE4 has been shown to have an important role in the regulation of vasopressin-induced translocation of AQP2 to the plasma membrane in cultured kidney cells (Stefan et al., 2007). In our study, AQP2 was detected in the human saccule together with two PDE4 members. Thus, it is possible that PDE4A and PDE4D control a cAMP pool involved in the regulation of AQP2 translocation. Functional studies indeed indicate an important role for AQP2 in the inner ear (Takeda et al., 2010), as is the case in the kidney. For example, it has been shown that expression of AQP2 is up-regulated in the cochlea and endolymphatic sac by the systemic application of vasopressin whereas it is down-regulated by systemic and local application of a vasopressin receptor 2 antagonist (OPC-31260; Takeda et al., 2010). The observation of high plasma vasopressin levels in Meniere’s disease suggests that components in the vasopressin–cAMP–AQP2 system, including cAMP PDEs, are promising targets for the development of therapies for diseases characterized by vertigo and hearing loss. However, although the vasopressin–cAMP–AQP2 system is believed to be important for water homeostasis in the inner ear, exactly what the role is for this system in the human saccule is not known and needs further investigation.

Two cGMP-degrading PDEs, PDE9A, and PDE10A, were shown to be expressed in the epithelium of human saccule. When it comes to the cGMP signaling system and the regulation of ion and water homeostasis in the inner ear much less is known as compared to the cAMP system. However, the cGMP-increasing hormone atrial natriuretic peptide has been suggested to regulate inner ear functions via the atrial natriuretic peptide receptor (Long et al., 2010), which is a membrane bound guanylate cyclase. Also, stimuli activating cGMP-dependent protein kinase

![Figure 4] Immunostaining of salt-inducible kinase (SIK) 1. SIK1 ([A] red), merged photos of SIK1 and DAPI ([B] blue). Lumen (lu), stroma (st), sensory epithelium (bar with two arrowheads). Bars 25 μm ([A],[B]).

![Figure 5] Immunostaining of the α-1 subunit of the N^+ , K^+ -ATPase. Merged photos of N^+ , K^+ -ATPase and DAPI ([A]), merged photos of N^+ , K^+ -ATPase and laminin ([B]) and laminin ([C]). Lumen (lu), stroma (st), basal membrane (bm), non-sensory epithelium (bar with one arrowhead ([A]). Bars 25 μm ([A]), 32.5 μm ([B],[C]).

![Figure 6] Hypothesis regarding how selected signaling components may interrelate in the inner ear to control ion and water homeostasis. PDEs and potential downstream targets of cAMP and cGMP signaling networks involving cAMP and cGMP activated kinases (PKA and PKG, not studied in this work), SIK1, the α-1 subunit of Na^+ , K^+ -ATPase, and AQP5 in the human saccule sensory epithelium are schematically represented. The presence of these signaling networks in the inner ear needs to be verified and studied at the functional level.
has been shown to cause translocation of AQP2 to the plasma
membrane in kidney epithelial cells (Bouley et al., 2000, 2005).
Furthermore, cGMP production is tightly linked to the nitric
oxide system, the soluble form of guanylate cyclase being acti-
vated by nitric oxide (Semaan et al., 2005). Excess nitric oxide
production is believed to mediate ototoxicity induced by cisplatin
and aminoglycosides by stimulating the production of reactive
oxygen species (Semaan et al., 2005; Hong et al., 2006; Rybak
et al., 2009). In addition, nitric oxide has been implicated in
mediating ototoxic effects associated with endolymphatic hydrops;
surgical induction of endolymphatic hydrops results in the upreg-
ulation of nitric oxide synthase II in cochlear and vestibular cells
(Semaan et al., 2005). Exactly what the role is for nitric oxide-
mediated induction of cGMP, or cGMP produced by soluble
 GUARDIAN CYCLES in response to other signals in the inner ear is not
known. Thus, the role for PDE9A and PDE10A in cGMP-related
processes needs further investigation. With regard to another
cGMP-degrading PDE, PDE5 (a cGMP-specific enzyme), verti-
tigo, and sudden hearing loss have been observed in patients
AQP9 DETERMINATION IN human saccule. To our knowl-
edge PDEs and SIK1 are new actors in the vestibular system and
the importance of these proteins in signaling networks regulat-
ing AQP4, Na+, K+-ATPase, and other components of relevance
for water and ion homeostasis remains to be established. It is
also important to verify the findings using other techniques than
immunohistochemistry since there is a potential for non specific
staining. Regarding the potential role in water and ion home-
ostasis for the components studied, this can be discussed in the
context of radial flow of endolymph. A number of studies chal-
lenge the largely accepted longitudinal flow theory which suggests
that endolymph is produced in the cochlear duct and flows in
an unidirectional pattern toward the endolymphatic sac, where
resorption occurs (Guild, 1927). The radial flow theory postu-
lates that endolymph is produced and absorbed throughout the
endolymphatic space. For example, using ionic markers Salt and
Ma (2001) provided evidence that control of endolymph home-
ostasis is distributed throughout the endolymphatic space. In
addition, a study using microscale analysis of proteins in the inner
ear tissues and fluids with emphasis on the endolymphatic sac,
otocia, and organ of corti support the existence of marked
regional differences in the mode and rate of endolymph protein
regulation implicating the role of various locally acting molecules
and ion channels in endolymph homeostasis (Thalmann et al.,
2006).

ACKNOWLEDGMENTS
The authors would like to acknowledge Gunnel Roos and Ann-
Kristin Holmén-Pålbrink for excellent technical assistance. The
authors would also like to acknowledge the staff at the surgi-
cal ward at the Department of Neurosurgery. This work was
supported by the Swedish Research Council Project 3362 (to
Eva Degerman) and 2611 (to Måns Magnusson); the Swedish
Diabetes Association; the Swedish Society of Medicine; the
A. Pålsson foundation and the Novo Nordisk Foundation,
Denmark.
identified as an abundant protein in the blood-labyrinth barrier that plays an essential role in the barrier integrity. PLoS ONE 6, e16547. doi: 10.1371/journal.pone.0016547


Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.