Role of aquaporins and regulation of secretory vesicle volume in cell secretion

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Received: August 16, 2007; Accepted: January 7, 2008

Abstract

In exocrine glands, secretory proteins synthesized in the rough endoplasmic reticulum (RER) exhibit vectorial transport from ER through a succession of membrane-bounded components such as Golgi complex, condensing vacuoles and secretory granules. The secretory granules migrate to particular locations within the cell close to the apical membrane prior to the release of their contents into the acinar lumen. Currently, to release intragranular contents, secretory granules have been demonstrated to transiently dock and fuse at ‘porosome’, a permanent cup-shaped structures at the cell membranes. Then swelling of secretory granules occurs to allow expulsion of intragranular contents. In this process, water and ion fluxes in the granule membrane appear to contribute to maintain secretory granule integrity and morphology via osmoregulation in secretory granules. Aquaporins (AQPs) are a family of small, hydrophobic, integral membrane proteins, which function as channels to permeate water and small solutes. The AQPs reside constitutively at the plasma membrane in most cell types. However, recent studies have demonstrated that the AQPs are present in secretory granules in exocrine glands, synaptic vesicles and intracellular vesicles in liver and kidney, implying that AQPs in secretory granules and vesicles are involved in their volume regulation. This paper reviews the possible role of AQPs on secretory granules, especially in exocrine glands, in secretory function.

Keywords: aquaporin • ion channel • granule swelling • secretory granules • porosome

Introduction

Aquaporins (AQPs) are a family of small, hydrophobic, integral membrane proteins of about 270 amino acids. In 1991, AQP1, formerly called CHIP28, was cloned as a first water channel and biophysically characterized [1, 2]. Sequence analysis of AQP1 demonstrated that AQP protein subunits consist of six α-helix transmembrane domains with inverted symmetry between the first three and last three domains ([3]; Fig. 1). The two loops between transmembrane helices 2-3 and 5-6 contain the signature amino acid sequence motifs of the AQPs, asparagine-proline-alanine (NPA) sequences. This predicted topology led to the ‘hourglass model’ of AQP structure, in which these two NPA-containing loops connect in the centre of the lipid bilayer and form a hydrophilic pore for water transfer through the lipid bilayer ([4]; Fig. 1). The hourglass of AQP structure has been confirmed using electron and X-ray crystal analysis of AQP1 [5, 6].

Biochemical and freeze fracture studies have indicated that AQP1 assembles in tetramer in the plasma membrane and each monomer contains a separate pore [4, 7]. The water selectivity of AQP1 has been suggested to be due to both the size-exclusion effect of the pore [8], and the orientation of asparagines in the NPA motifs providing necessary hydrogen-bonding interactions to isolate water molecule and avoid the passage of protons through the pore.
Secretory granule or vesicle is formed from the condensing membrane-bounded compartments including the Golgi complex. Proteins are vectorially transported from rough endoplasmic reticulum (RER) through a succession of synthesized in the Golgi complex, then to secretory granules or vesicles for their volume regulation.

In various exocrine and endocrine cells, secretory proteins are synthesized in rough endoplasmic reticulum (RER). They are vectorially transported from RER through a succession of membrane-bounded compartments including the Golgi complex, condensing vacuoles, and secretory granules and vesicles. The secretory granule or vesicle is formed by the condensing vacuole, which buds off the trans face of the Golgi complex. In the condensing vacuole, secretory proteins are present as dilute form. In a subsequent packing process, the proteins are condensed. During this process, it is conceivable that transport of ions and water through secretory granule membrane is necessary for the protein condensation, although there is not yet complete agreement on the basic principles involved.

The secretory granules and vesicles move to particular locations within the cell close to the plasma membrane prior to the release of their contents to the outside. Since the secretory vesicles have been suggested to fuse with the plasma membrane by using electron microscopy [31], it has been considered that cell secretion involves the fusion of membrane-bound granules and vesicles at the plasma membrane at the secretory sites, and the release of intragranular and intravesicular contents to the extracellular sites. Currently, by using atomic force microscopy, it has been demonstrated that secretory granules and vesicles transiently dock and fuse at ‘porosome’, a permanent cup-shaped structure at the cell membrane, and swell to allow expulsion of their contents [32-36]. Therefore, secretion involves the fusion of the granule or vesicle membrane with porosome in the plasma membrane, followed by the release of their contents outside. Secretion without stimulation is referred to as ‘constitutive secretion’, but cell secretion following a stimulus is termed ‘regulatory secretion’.

During the secretory process, regulation of secretory granule or vesicle volume is important, in which contribution of various ion channels has been demonstrated [21, 26, 37–39]. This paper reviews the possible role of AQPs on secretory granules and vesicles in secretory function, especially in exocrine glands.

**Role of AQPs in cell secretion**

AQP5 is a water-selective channel protein widely expressed in exocrine glands [40–42]. In rat duodenal Brunner’s gland, vasoactive intestinal polypeptide (VIP) has been reported to increase the flow rate as well as bicarbonate and protein output [43]. Parvin et al. [23] have demonstrated that AQP5 localizes in the secretory granule membrane and the apical membrane in rat Brunner’s gland by immunohistochemistry and electron microscopic immunohistochemistry, and that the AQP5 level in the apical membrane is increased by VIP stimulation. This observation suggests that the AQP5 translocates from the secretory granule membrane to the apical membrane on secretion provoked by VIP. In the rat parotid gland, AQP5 is localized in the apical membrane including the intracellular canaliculi of acinar cells [41]. In rat parotid acinar cells, AQP5 has also been reported to translocate from the intracellular vesicles to the apical membrane in vitro in response to stimulation with muscarinic agonists, which induce water secretion and feeble release of the digestive enzyme amylase [44]. However, the immunofluorescence and immunoelectron microscopic studies demonstrated that AQP5 was predominantly localized in the apical plasma membrane in the mouse parotid and submandibular glands after stimulation or inhibition of secretion in vivo.
indicating that no changes in the subcellular localization of AQP5 occurs [45]. On the other hand, it has been reported that immunostained AQP5 was scattered as clusters in the submembranous area of the acinar cells in the rat injected with the β-agonist isoproterenol, which provokes amylase secretion in parotid acinar cells [42]. In the mouse parotid gland, AQP5 was localized along the apical membrane and its small invaginations formed by fusion of secretory granules after isoproterenol-administration [46]. The fluidity of the primary secretion has been proposed to be important for the release of intragranular contents from membrane-bound secretory granules in pancreas [47] and parotid gland [48]. It is most likely that components of granule membrane such as ion channels might insert into the apical membrane during fusion process at porosome, and subsequently salt and water would flush out the stored macromolecules into the acinar lumen and provide for an appropriate amount of fluid to be secreted with the proteins [47, 48].

Individual secretory events can be visualized with sulforhodamine B (SRB), a fluorescent fluid-phase polar tracer, as the formation of docked-granule profiles at apical membrane using two-photon excitation microscopy, since SRB remaining in the luminal region rapidly diffuses into granules fused with the plasma membrane [49]. When secretory events in rat parotid acini were investigated using the two-photon microscopy, the β-agonist isoproterenol provoked cell secretion, docked-granule profiles, at apical region as Figure 2 shows (Sugiya, Nemoto & Kasai, in preparation). However, the profiles formed disappeared soon. This observation suggests that SRB diffused into granules fused with the plasma membrane may subsequently be diluted by water rapidly, and implies function of AQP5 in secretory granule membrane.

Role of AQPs in secretory granule swelling

In cell secretion process, role of secretory granule swelling has been investigated. In beige mouse mast cells, membrane fusion has been demonstrated to precede secretory granule swelling during cell secretion by the studies with electrophysiological membrane capacitance measurements [50, 51], which proved that osmotic swelling is not required for fusion. It has been inferred that secretory granule swelling is necessary to stabilize and widen the fusion pore and is caused by movement small solutes through the fusion pore into the granule matrix.

On the contrary, secretory granule swelling has been proposed to be prerequisite for secretory granule fusion with plasma membrane [52–56]. On the hypothesis, it has been considered that swelling of secretory granules results in a build-up of pressure for allowing expulsion of intragranular contents and the extent of secretory granule swelling dictates the amount of intragranular contents expelled during secretion.

In rat pancreas, AQP1 had been demonstrated to be localized at zymogen granules, the membrane-bound secretory vesicles...
The amino and carboxyl domains of AQP1 have suggested to localize at the luminal side of zymogen granule membrane, because immunoreacted signal of AQP1 was detected by immunoblot analysis when anti-AQP1 antibody to the carboxyl terminus of AQP1 was introduced into to the zymogen granules permeabilized by streptolysin-O, no immunoreacted signal was detectable in the sample of intact zymogen granules pre-exposed to the anti-AQP1 antibody, and immunogold-labelling was observed at the inner side of membrane of streptolysin-O-permeabilized zymogen granules on the immunoelectron microscopy using the anti-AQP1 antibody [21]. This topology is curious, because the hydropathy analysis suggests that AQPs have six putative helical domain, and the studies utilizing epitope tagging, AQP-reporter chimeras, and site-specific antibodies confirmed this fundamentally topological organization and indicated that the amino and carboxyl termini are cytoplasmically oriented [57].

The contribution of AQP1 to zymogen granule swelling was studied [21]. Fusion of zymogen granules with plasma membrane has previously been reported to be facilitated by activation of the trimeric GTP-binding protein G\textsubscript{i}, in zymogen granule membrane by \textit{in vitro} fusion assay [58, 59]. Subsequently, the guanosine 5'-triphosphate (GTP)-binding proteins has been shown to contribute to secretory granule swelling using atomic force microscopy [37], which provides three-dimensional data with the structure and dynamics of single biomolecule, living cells and organelles including secretory granules [60]. These observations suggest that zymogen granule swelling is an important prerequisite for zymogen granule fusion with plasma membrane. Cho et al. [21] have been demonstrated that the GTP-mediated increase in granule volume and water entry into granules determined using atomic force microscopy and tritiated water, respectively, were inhibited by \textbf{HgCl}2, an inhibitor of AQP1 [21], or by insertion of anti-AQP1 antibody into the zymogen granules. Therefore, it is most likely that AQP1 contributes to zymogen granule fusion with plasma membrane and expulsion of granule contents during secretion in pancreatic acinar cells.

In synaptosomes and synaptic vesicles from rat brain, Kelly et al. [39] examined whether synaptic vesicle swelling is similarly regulated to the zymogen granules in exocrine pancreas using atomic force microscopy. Since size of synaptic vesicles increased in the presence of GTP and the Go/Gi stimulator mastoparan, an amphiphilic tetradecapeptide from wasp venom, synaptic vesicle swelling \textit{via} activation of heterotrimeric GTP-binding protein has been demonstrated [39]. Subsequently, water channels, AQPs1 and AQP6, associated with synaptic vesicles isolated from rat brain were detected [26]. Since Go\textsubscript{3}, was also detected as the major heterotrimeric GTP-binding protein in synaptic vesicles, Jeremic et al. [26] examined the role of AQPs in synaptic vesicle swelling \textit{via} the activation of Go. The effects on synaptic vesicle size of Go protein stimulators, GTP and mastoparan, were examined using various approaches, such as photon-correlation spectroscopy, right-angle light scattering and atomic force microscopy. Consequently, swelling of synaptic vesicles rapidly occurred in response to GTP and mastoparan. However, in the presence of the AQP inhibitor \textbf{HgCl}2, the stimulatory effect of mastoparan on synaptic vesicle swelling was significantly abrogated [26]. These observations strongly suggest that synaptic vesicle swelling is caused by a \textit{Go}-regulated, AQP-mediated water entry in secretory vesicles and involved in neurotransmitter expulsion, as suggested in pancreatic zymogen granules.

In secretory granules isolated from the rat salivary parotid gland, AQPS has been demonstrated to localize on the membrane of secretory granules by immunoblot analysis and immunoelectron microscopy [22]. On the immunoelectron microscopy using anti-AQPS antibody to the carboxyl terminus of AQPS, the immunogold particles were detected at the outside of secretory granule membrane. To study the function of AQPS in the parotid secretory granules, Matsuki et al. [22] utilized a quantitative \textit{in vitro} assay involving rapid osmotic swelling and end-point measurements of granular osmotic lysis, which has been used for the investigation of ion conductance in secretory granules [61]. In this assay system, it has been demonstrated that anti-AQPS antibody induces secretory granule swelling and lysis in iso-osmotic KCl solution, suggesting that inhibition of AQPS function causes secretory granule swelling and lysis. In secretory granules of the rat parotid gland, expression of the heterotrimeric GTP-binding protein G\textsubscript{o} and G\textsubscript{o}-regulated Cl– conductance have been reported [62, 63]. However, AQPS regulation by GTP-binding proteins in the secretory granule is still unknown.

**Role of AQPs in ion permeation**

On the basis of studies using atomic force microscopy in pancreatic acinar cells and isolated zymogen granules, Jena et al. [37] proposed that K\textsuperscript+ and Cl\textsuperscript– channels in the granule membrane need to induce granule swelling during secretion to prevent collapse of zymogen granules. Ion fluxes through K\textsuperscript+ and Cl\textsuperscript– channels in the granule membrane and osmotic swelling thus appear to contribute to maintain granule integrity and morphology in secretory function.

When the molecular mechanism of swelling of pancreatic zymogen granules was studied, detergent-solubilized zymogen granules co-isolate the inwardly rectifying K\textsuperscript+ channel IRK-8 and the chloride channel CLC-2, in addition to other proteins such as AQPS, G\textsubscript{o}3 and phospholipase A\textsubscript{2} after immunoprecipitation with a monoclonal AQPS antibody [64]. Exposure of zymogen granules to either the K\textsuperscript+ -channel blocker glyburide or the phospholipase A\textsubscript{2} inhibitor ONO-RS-082 blocked GTP-induced zymogen granule swelling. In liposomes reconstituted with the AQPS immunoisolated complex from solubilized zymogen granules, swelling in response to GTP occurred, but the GTP effect was abolished by glyburide or ONO-RS-082. In the planar lipid membranes reconstituted with the immunoisolate complex, conductance corresponding to the passage of K\textsuperscript+ was decreased by glyburide or an anti-AQPS antibody. These observations suggest that G\textsubscript{o}3-phospholipase A\textsubscript{2} -mediated pathway and K\textsuperscript+ channels are involved in AQPS regulation [64]. It has been reported that AQPS expressed in \textit{Xenopus} oocytes conducted cations [65] and the water permeability function was inhibited by the K\textsuperscript+ channel.
blocker, tetraethyl ammonium [66]. When AQP1 purified from Saccharomyces was reconstituted into planar lipid bilayers, cyclic GMP-induced ion conductance was detected, although the ion channel number was exceedingly low [67]. The amount of similarity between K+ channels and AQP1 has been demonstrated [68]. These observations imply that AQP1 is involved in K+ flux in zymogen granules directly or indirectly, although crystallography studies have been demonstrated that AQP1 is a water-specific channel [6].

Matsuki et al. [22] have also demonstrated that the anti-AQP5 antibody-induced granule lysis is inhibited in the absence of Cl− or in the presence of 4,4-diisothiocyanostilbene-2,2'-disulfonic acid, an anion channel blocker in the reaction mixture. There is no evidence that AQP5 acts as ion channels in the AQP5-expressing oocytes [65, 69]. The presence of Cl− conductance in the secretory granule membrane of the rat parotid gland has been demonstrated [48, 70]. In airway secretory glands, the expression of cystic fibrosis transmembrane conductance regulator (CFTR), a cAMP-activated Cl− channel, has been reported [71], and CFTR has been suggested to be related to the ion content in the secretory granules and granule expansion [72]. As a consequence, we hypothesize that a balance of water permeation via AQP5 and Cl− conductance is necessary for secretory granule volume regulation. Currently, the hypothesis that AQPs function as osmotic and turgor sensors rather than water channels, the sensor hypothesis, has been advocated [73, 74]. Therefore, AQP5 appears to act as an osmotic sensor in the secretory granules of the parotid gland, although further studies with precise mechanisms of the relationship between AQP5 and the Cl− channels are necessary.

**Fig. 3.** Localization of AQP6 in rat parotid secretory granule membrane. Protein expression of AQP6 in parotid secretory granule membrane (granule) was detected by western blotting using anti-AQP6 antibody. In granule membrane, 33 kDa band of AQP6 was detected. Kidney medul- lae epithelial cells (kidney) were used as a positive control.

**Fig. 4.** Localization of AQP6 in rat parotid secretory granules. Immunoelectron microscopy in ultrathin cryosection was carried out. Anti-AQP6 antibody was labelled with 10 nm colloidal gold-conjugated secondary antibody. (A) Colloidal gold particles were observed in parotid secretory granules as indicated (solid arrowheads) and in other spots (open arrowheads). (B) Colloidal gold particles were observed both in outer and inner leaflets of secretory granule membrane. SG, secretory granule; Lum, lumen; Mv, microvilli; ICS, intercellular canalicu- lus. Bar, 0.5 µm.
The function of most AQPs is well known to be inhibited by mercurial agents which bind the sulfhydryl (SH) group of cysteines. In AQP5, a cysteine at residue 182 in loop E, a hydrophobic loop (Fig. 1), is considered to be a mercury-sensitive domain [75], and corresponds to the known mercurial-inhibitory site [76]. Therefore, we examined the effect of HgCl$_2$ on lysis of secretory granule in the parotid gland. Interestingly, HgCl$_2$ clearly induced parotid secretory granule lysis [77]. The HgCl$_2$-induced granule lysis was also completely blocked in the presence of 8-mercaptoethanol, a protective agent for SH groups, which has been demonstrated to restrain the effect of HgCl$_2$ on AQP5 [78]. These observations were thought to support the view that the inhibition of AQP5 function causes secretory granule lysis.

However, it has been demonstrated that AQP6 function is activated by HgCl$_2$ [79, 80]. AQP6 genes have been identified in rat and human kidneys with high homologies to AQP0 and AQP2 [81, 82]. Subsequently, AQP6 has been found to be localized in intercellular sites of acid-secreting $\alpha$-intercalated cells from renal-collecting duct [27]. It has been considered that AQP6 functions not as a water channel but as an anion channel, because permeation by anions in response to acidic pH or Hg$^{2+}$ activation in Xenopus Laevis oocytes expressing AQP6 was found [79] and Hg$^{2+}$-activated ion conductance was verified by single-channel recordings of the oocytes [83]. In Xenopus Laevis oocytes expressing AQP6, Hg$^{2+}$ has also been demonstrated to stimulate transport of glycerol and urea [80]. Currently, we found the presence of AQP6 associated with secretory granule membranes in the rat parotid acinar cells [84]. In the granule membrane isolated from the purified secretory granules, anti-AQP6 antibody specifically recognized 33 kDa band by western blotting, indicating the presence of AQP6 in secretory granule membrane of rat parotid acinar cells, as Figure 3 shows. To confirm the presence of AQP6 in parotid secretory granules, immunoelectron microscopy in ultrathin cryosection was carried out (Fig. 4). Close observation of immunogold labelling of secretory granules demonstrates gold labelling both the inner as well as outer leaflet of the granule membrane. In HEK239 cells, when a green fluorescence protein (GFP) tag was added to the N-terminus of AQP6, GFP-AQP6 was redirected to the plasma membranes [85]. In HEK239 cells expressing GFP-tagged AQP6, AQP6 has also been demonstrated to function as an anion channel with the halide permeability sequence: NO$_3^->I^->Br^->Cl^->F^-$ [85], strongly supporting that AQP6 functions as anion channel in mammalian cells. Therefore, it is most likely that AQP6 functions as a Cl$^-$ channel and contributes to regulation of osmoregulation in secretory granules of the parotid gland, although further studies need to elucidate AQP6 function in secretory granule. In synaptic vesicles, HgCl$_2$ inhibited the GTP-induced vesicle swelling, despite of the presence of AQP6 [26]. Therefore, role of AQP6 in the synaptic vesicle swelling is unknown.

**Conclusions**

Secretion of macromolecules in exocrine and endocrine cells occurs through docking and fusion of secretory granule membrane at poresome in plasma membrane and the subsequent discharge of secretory granule contents [32–36]. It has been suggested that secretory granule swelling is involved in the exocytotic process. A specific set of ion channels in secretory granule membrane has been proposed to contribute to the secretory granule swelling [21, 26, 37–39]. AQPs in secretory granules membrane have to be demonstrated to be concerned in the granule swelling and may contribute to release of contents in secretory granules, which could provide a new information with AQP functions. In pancreatic zymogen granules, functional relationship between AQP1 and K$^+$ channel has been demonstrated [64]. In parotid secretory granules, the presence of AQP5 and AQP6, which are water-selective and Cl$^-$-permeable channels, respectively, has been demonstrated [22, 84]. Therefore, the relationship between water channels and ion channels has to be elucidated. To elucidate the function of AQPs in secretory granules, specific, non-toxic AQP inhibitors and specific AQP antibodies will be useful.

Cell secretion can be greatly accelerated following an appropriate cellular signal, which is called ‘regulatory secretion’. Regulatory secretion is dependent on intracellular Ca$^{2+}$ or other intracellular signals. AQP4 function has been reported to be regulated by protein kinase C [86]. Studies with such a regulation on AQPs in secretory granules may be important to elucidate the functions.

**Acknowledgements**

This study was supported in part by a Nihon University Multidisciplinary Research Grant for 2006-2007 and a Grant of Oral Health Science Center HRC7 from Tokyo Dental College.

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