

Structural basis for conductance by the archaeal aquaporin AqpM at 1.68 Å

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To explore the structural basis of the unique selectivity spectrum and conductance of the transmembrane channel protein AqpM from the archaeon *Methanothermobacter marburgensis*, we determined the structure of AqpM to 1.68-Å resolution by x-ray crystallography. The structure establishes AqpM as being in a unique subdivision between the two major subdivisions of aquaporins, the water-selective aquaporins, and the water-plus-glycerol-conducting aquaglyceroporins. In AqpM, isoleucine replaces a key histidine residue found in the lumen of water channels, which becomes a glycine residue in aquaglyceroporins. As a result of this and other side-chain substituents in the walls of the channel, the channel is intermediate in size and exhibits differentially tuned electrostatics when compared with the other subfamilies.

integral membrane protein | x-ray crystallography | water channel | gas

Aquaporins are a large family of transmembrane channel proteins that facilitate the passive, but highly selective, movement of water, small neutral alditols, including glycerol, or CO₂ across cell membranes (1). These channels are found throughout all domains of life and are fundamental elements in the osmoregulation of many organisms. The selectivity and biophysical characteristics of several aquaporins have been described through atomic resolution structures. The atomic resolution structures of aquaporins GlpF (2), Aqp1 (3, 4), AqpZ (5), Aqp0 (6, 7), and now most recently, AqpM, show high similarity in their tertiary structures. However, despite the similarity in topology and quaternary structure across the family, *in vivo* and *in vitro* water and glycerol conductance assays (8–10) have shown that each member exhibits unique conductance rates and displays selectivity spectra that extend to the selective permeation of small neutral molecules that include glycerol, urea, and CO₂.

The structural congruity that spans this family points to the side-chain variations in the channels as the main basis for the divergence in the selectivity and conductance rates between different aquaporins. For example, the crystal structures revealed the integral role of the extended loop that leads into the central plane of the membrane bilayer from each side of the membrane to the conserved -NPA- motifs that initiate the half-membrane spanning helices. These features are a signature for the entire family. Each of these loops provides a ladder of four carbonyl oxygens from successive amino acids in the sequence that are presented into the lumen of the channel. Forming a helical set of hydrogen-bond acceptors from one side to the other along the axis of the channel, they generate a ladder of hydrogen-bond-accepting groups spaced ≈ 2.8 Å apart in distance through the channel. They determine a key element of the chemical environment for the coordination and conductance of the permeants through the lumen of the channel. Also, the side chains that form the selectivity filter influence the electrostatics and diameter of the channel at its narrowest juncture (2) and separate the water-selective aquaporins such as Aqp1 and AqpZ from the glycerol-conducting subfamily (aquaglyceropor-

ins) that includes the *Escherichia coli* glycerol channel GlpF and human aquaporins 3, 7, 9, and 10.

The discovery and characterization of AqpM (11) from the methanogenic archaeon *Methanothermobacter marburgensis* as an aquaporin confirmed the presence of aquaporins in archaea (12). Because of the absence of any other aquaporins in the genome of *Methanothermobacter marburgensis*, the primary function of AqpM is most likely to facilitate the movement of water across the plasma membrane in response to osmotic gradients of its environment. *In vitro* and *in vivo* conductance assays verified that AqpM is indeed a water-conducting channel (13), albeit with a lower permeability rate for water (P_f) than other water-selective aquaporins such as AqpZ [$P_f(\text{AqpM}) = 57 \pm 4 \mu\text{m/s}$ (13); $P_f(\text{AqpZ}) = 330 \mu\text{m/s}$ (8)]. AqpM has also been observed to conduct glycerol at a low rate, but in *Methanothermobacter marburgensis*, which relies on CO₂ as its sole carbon source, this function may prove to be biologically irrelevant (13).

Because of the high similarity in the structures of most aquaporins, several unusual qualities of AqpM provide awareness for deciphering the mechanism of selectivity and conductance by aquaporins. First, an isoleucine residue replaces the highly conserved histidine residue located in the selectivity filter of water-selective aquaporins. This difference may represent an adaptation specific for function in addition to the conduction of water. In support of this speculation, initial studies indicate the conductance of CO₂ by AqpM (Y.K., unpublished data). The identification of certain aquaporins as CO₂ channels has been a topic of some interest and controversy (14, 15), although recent studies have shown the conductance of CO₂ by bovine Aqp1 (16) and tobacco aquaporin NtAQP1 (10). Notably, these two aquaporins are highly expressed in membranes involved in the high traffic of CO₂, namely the membranes of red blood cells and plant leaves. The demand for specific and efficient conductance of this gas by certain cells and the characterization of aquaporins with this functional capacity supports the possibility that aquaporins may be this elusive gas channel.

Here, we report the atomic structure of AqpM to 1.68-Å resolution. Based on this structure, we elaborate the relationships between the structure, rate of conductance, and selectivity by this transmembrane channel protein for water and glycerol. The structure of AqpM supports the role of the channel vestibules in preselecting the permeants and beginning the energetically unfavorable process of stripping the hydration shell from around the molecule before it is transported into and through the narrow portion of the channel.

Conflict of interest statement: No conflicts declared.

Abbreviation: OG, octyl- β -D-glucoside.

Data deposition: The atomic coordinates have been deposited in the Protein Data Bank, www.pdb.org (PDB ID codes 2F2B and 2EVU).

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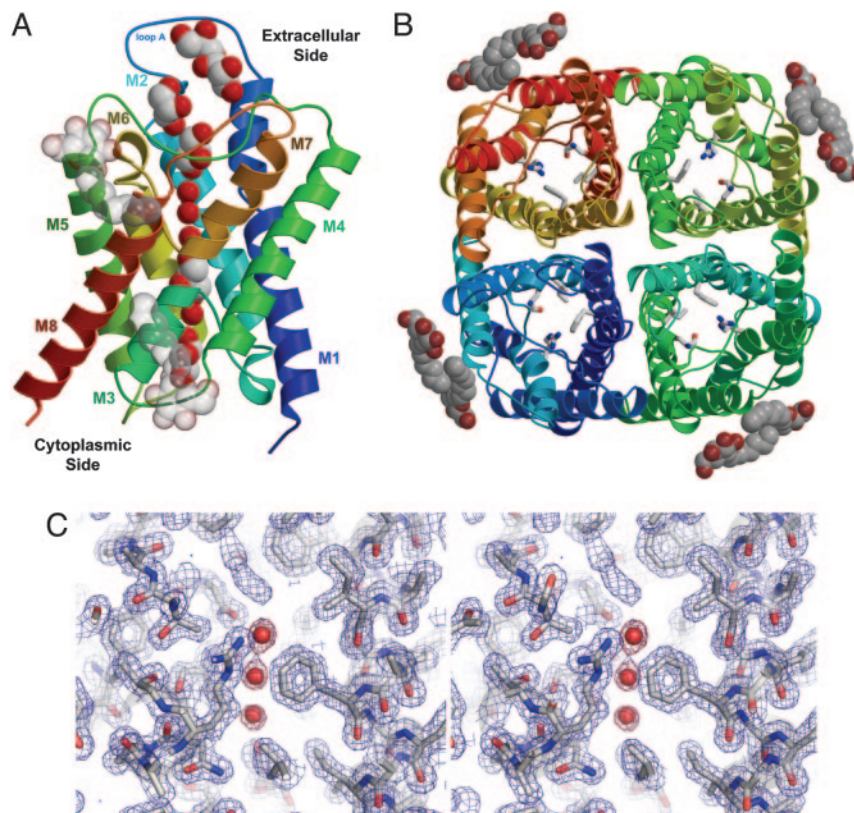


Fig. 1. Monomer and tetramer view of AqpM. (A) AqpM monomer viewed parallel to the plane of the membrane. The six transmembrane (M1–M2, M4–M6, and M8) and two half-membrane (M3 and M7) spanning helices are labeled M1–M8. The glycerol and water molecules found in the channel are represented as spheres. The two OG detergent molecules identified in the structure are represented as transparent spheres. The channel is readily identifiable by the line of water and glycerol molecules coordinated inside the channel. (B) The AqpM tetramer viewed down the 4-fold axis of the tetramer from the extracellular aspect of the membrane. Four residues that form the selectivity filter are shown in stick rendering. (C) Stereoview of the sigma-weighted $2F_o - F_c$ electron density at the selectivity filter. The map is contoured at 1.2σ . The figure was made with MOLSCRIPT (25), RASTER3D (26), and PYMOL (27).

its equivalent in other aquaporins. This long loop provides a structural role in forming a part of the tetramer interface. It also folds to generate a hydrophobic pocket on the extracellular surface that is bound by glycerol and water molecules and is a part of the channel vestibule.

The differences between the low-resolution (2.3 Å) and high-resolution (1.68 Å) structures are minimal with a rms deviation of 0.17 Å between the two. However, differences exist in the location and presence of water, glycerol, and detergent molecules associated with the protein. The detergent molecules are only in the low-resolution structure. The location of water and glycerol molecules in the channel are also slightly different. The pore of the high-resolution structure is occupied almost exclusively by water molecules, with one glycerol molecule present at the -NPA- region of the channel. The low-resolution structure reveals four water molecules (HOH1–HOH4) and three glycerol molecules (G1–G3) lined up in single file along the channel, with each molecule hydrogen-bonded to its neighbor, while the main-chain carbonyl oxygens that line the channel provide hydrogen-bond acceptors for the permeant molecules (Fig. 2). Traveling from the extracellular side of the channel to the cytoplasmic side down the transmembrane axis of the channel, a glycerol (G1) molecule occupies the first position, at the mouth of the channel, stabilized by a hydrogen bond (2.67 Å) between the hydroxyl group of C2-OH and the main-chain carbonyl oxygen of G133(135) (all AqpM residue numbers are followed by GlpF numbering in parenthesis for reference). G1 is followed by three successive water molecules (HOH1, HOH2, and HOH3), located adjacent to the carbonyl oxygens of G195(198) (2.81 Å),

S196(199) (2.68 Å), and S197(200) (2.56 Å), respectively. HOH3 is followed by G2, which is adjacent to both the aquaporin canonical NPA motifs of helices M3 and M7. G2 is followed by

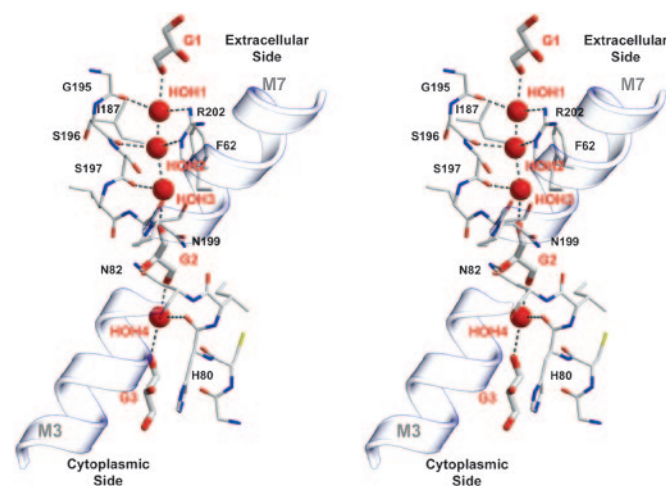


Fig. 2. AqpM channel architecture. Stereoview of the channel with potential hydrogen bonds to the waters (HOH1–H4) and glycerols (G1–G3) shown in gray dotted lines. Helices of M3 and M7 are rendered in cartoon representation, while the loop portion residues are represented as sticks. The main-chain carbonyl oxygens of the two loops form the conserved ladder-like structure of hydrogen-bond acceptors that line the channel. N82 and N199 are the asparagines that make up the canonical aquaporin NPA motifs.

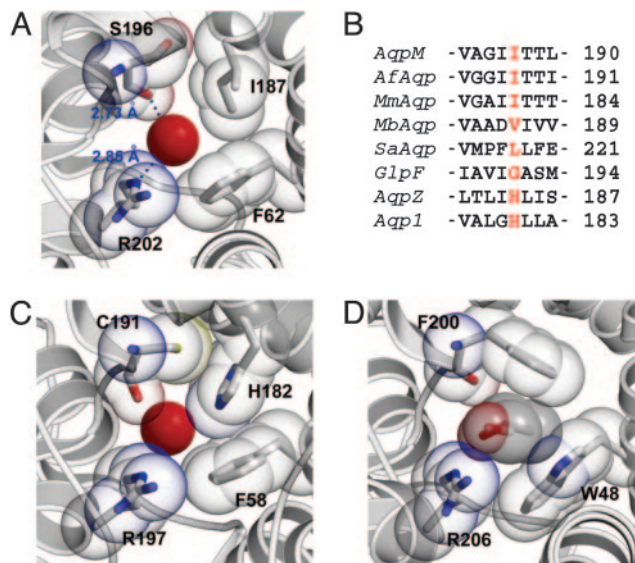


Fig. 3. Aquaporin selectivity filters. (A) The residues of the selectivity filter of AqpM (R202, F62, S196, and I187) are depicted with the water molecule (HOH2) coordinated at the center of the filter. The potential hydrogen bonds from HOH2 to S196 and R202 are depicted as a blue dotted line. (B) The sequence alignment of six aquaporins. I187 from AqpM and corresponding residues in the other aquaporins are highlighted in the red box. In water-selective aquaporins, represented here by Aqp1 and AqpZ, a conserved histidine residue occupies this position. In aquaglyceroporins such as GlpF, a glycine residue occupies this position. Similar to AqpM, the two other archaeal aquaporins [*A. fulgidus* aquaporin (AfAqp) and *Methanosarcina barkeri* aquaporin (MbAqp)] possess aliphatic residues at this position. SaAqp, *S. acidocaldarius* aquaporin. MmAqp, *Methanococcus maripaludis* aquaporin. (C and D) The selectivity filters of Aqp1 (C) and GlpF (D) highlighting the similarity and differences between an aquaporin and aquaglyceroporin. A glycerol molecule is seen only in the selectivity filter of GlpF (D).

HOH4, which is stabilized by a hydrogen bond to the carbonyl oxygen of H80 (3.10 Å). Lastly, G3 occupies the cytoplasmic end of the channel.

The channel constricts to its narrowest point at the selectivity filter. The filter is formed by the side chains of R202(206), F62(48), and I187(191) and the carbonyl oxygen of S196(191) (Fig. 3A). The N ϵ of the guanidino group of R202(206) along with the carbonyl oxygen of S196(191) forms the hydrogen-bond donor and acceptor pair along half of the roughly circular filter (when viewed normal to the membrane plane), while the hydrophobic residues F62(48) and I187(191) form the other half. The topology of the selectivity filter is almost identical to that of other water-selective aquaporins, with the exception of I187(191), which replaces the highly conserved histidine residue (Fig. 3B). The arrangement of these residues constricts the diameter of the channel [all channel diameters cited were as calculated by the HOLE2 (20) program] to its minimum of 2.54 Å, 0.68 Å wider than that of the equivalent site in the water-specific Aqp1 (1.86 Å) (Fig. 3C) and 0.60 Å narrower than that of the aquaglyceroporin GlpF (3.14 Å) (Fig. 3D). Notably, this calculated diameter is similar to that of the diameter of a water molecule (2.8 Å) and presents good accommodation for the passage of water. Although the diameter calculation represents the largest spherical molecule that can pass through a roughly circular pore, this value presents a fairly good approximation even for the noncircular aquaporin channel because it represents the largest sphere that can pass through this pore.

Three highly ordered water molecules (HOH1–HOH13) are found at or near the selectivity filter in both the high- and low-resolution structures. HOH1 sits on the extracellular side of

the selectivity filter and is hydrogen-bonded to the carbonyl oxygen of G195(190) (2.73 Å) and NH2 of R202(206) (2.85 Å). HOH2 is coordinated at the cytoplasmic side of the selectivity filter and is hydrogen-bonded to the N ϵ of R202(206) (2.89 Å) and the carbonyl oxygen of S196(191) (2.72 Å). HOH3 is located adjacent to HOH2 toward the cytoplasmic side and is hydrogen-bonded to the carbonyl oxygen of S197(192) (2.57 Å). The three water molecules are also hydrogen-bonded to each other, with HOH1 and HOH2 hydrogen-bonded to each other (2.5 Å), as are HOH2 and HOH3 (3.0 Å) (Fig. 2).

Proteoliposome Swelling Assay. After cleavage of the affinity tag, assays using purified AqpM reconstituted into liposomes reveal that it is active and functional in conductance of water (Fig. 4A). The curves represent the measured rate of vesicular shrinkage as a result of water efflux out of the proteoliposomes. This efflux is induced when vesicles are exposed to a sharp sucrose gradient (570 mM sucrose) in a stopped-flow light scattering measurement. The curves are fitted with a single exponential function as described by Borgnia and Agre (9), and the results indicate water permeability (P_f) of $60.5 \pm 3 \mu\text{m/s}$, comparable to that observed by Kozono *et al.* (13) using samples with the affinity tag still present on the samples. Our results show that the presence or absence of the affinity tag does not affect the water-conductance activity of AqpM.

Discussion

Water Conductance and Channel Architecture. The presence of only water molecules at the selectivity filter correlates with the biophysical characterization of AqpM with preferential selection for water over glycerol. The two water molecules coordinated at the selectivity filter (HOH1 and HOH2) are both hydrogen-bonded to R202(206) and the main-chain carbonyl oxygens of S196(191) and G195(190), respectively (Fig. 2). When visualized in space filling representation (Fig. 3A), the selectivity filter is wide enough to accommodate the passage of water and highly restrictive toward the passage of glycerol. With some movement of the side chains, the channel may expand enough to allow the passage of glycerol, but the channel seen in this structure reveals why the rate of the passage of glycerol is very low when compared with the rate seen for the glycerol transporter GlpF. The high concentration of glycerol (10% vol/vol) in the crystallization condition accounts for its presence in the channel, but of all of the crystal structures of aquaporins (GlpF, Aqp1, AqpZ, and Aqp0), only that of the aquaglyceroporin GlpF has a glycerol molecule present in the selectivity filter (Fig. 3D). Also, despite the slight differences in the location of water and glycerol molecules in the channel between the high- and low-resolution structures, the location of two water molecules at the selectivity filter are identical between the two structures reported here. Therefore, the coordination of the water molecule at the selectivity filter, despite the presence of glycerol molecules at other regions of the channel, and the restrictive diameter of the channel represents a steric and electrostatic environment that is clearly adapted for the stabilization of water or chemically similar molecules at this site.

Based on amino acid sequence variations, structures, and permeation studies, the aquaporins can be classified into two distinct subfamilies, water-selective and glycerol-selective. The structure and sequence of AqpM positions it in a unique place between these two classes and establishes a paradigm applicable for the archaeal aquaporins. The substitution of the highly conserved histidine residue [H182(191) in bAQP1, H174(191) in AqpZ] in the selectivity filter of AqpM with an aliphatic I187(191) is a feature seen only in the archaeal aquaporins [I188(191) in *Archaeoglobales fulgidus* aquaporin, V186(191) in *Methanosarcina barkeri* aquaporin, I181(191) in *Methanococcus maripaludis*, and L218(191) in *Sulfolobus acidocaldarius*] (Fig.

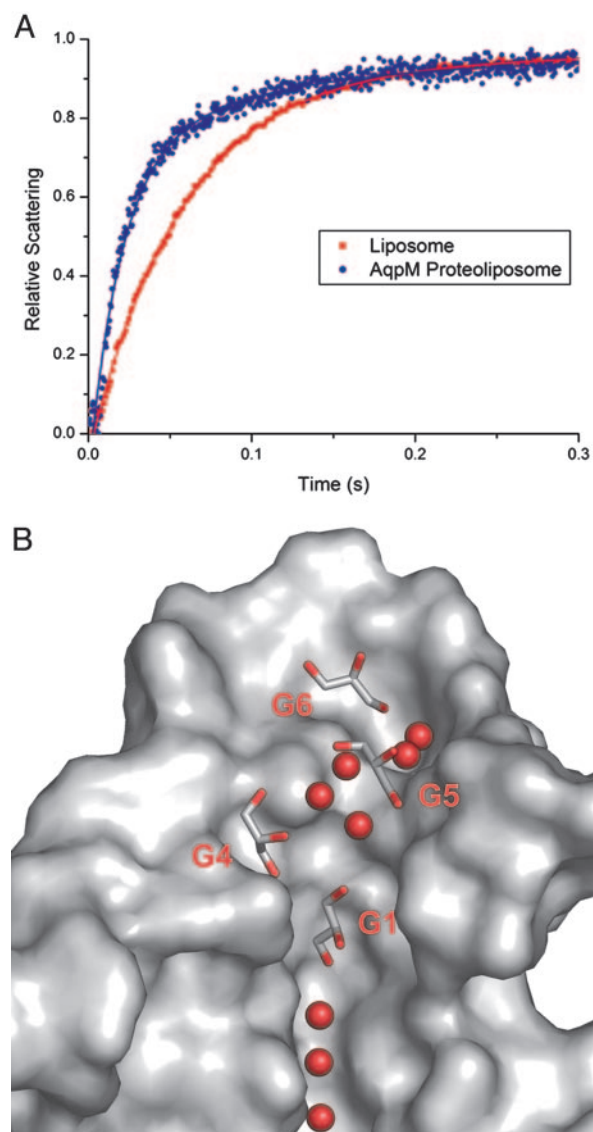


Fig. 4. Water conductance by AqpM and water and glycerol in the vestibule of the channel. (A) Water permeability of liposome-reconstituted AqpM. Liposomes reconstituted with purified AqpM or control liposomes were mixed at 12.2°C with an equal volume of hyperosmolar solution (570 mM sucrose). The reduction in vesicular volume caused by water efflux, resulting in an increase in light scattering, was observed in a stopped-flow apparatus for 0.5 s. The data were normalized between zero and unity and fitted to an exponential rise to the maximal value curve. Osmotic water permeabilities constants were calculated as follows: $P_f(\text{liposomes}) = 15.5 \pm 0.7 \mu\text{m/s}$, $P_f(\text{AqpM proteoliposomes}) = 60.5 \pm 3 \mu\text{m/s}$. (B) Shown is the surface rendering of AqpM with helices 3, 4, 7, and 8 removed to highlight the channel and vestibule. Three glycerol (G4–G6) and five water molecules can be seen in the pocket of the vestibule on the extracellular side of the channel. G4 is approximately in the optimal orientation before entering the narrow portion of the channel.

3B). Although this is only a single residue difference, this histidine residue is a highly conserved feature of water-selective aquaporins and represents one of the key distinctions between the two major subfamilies. In other respects, the selectivity filter of AqpM resembles that of the water-selective aquaporins. Two of the four residues at the selectivity filter of AqpM, R202(206) and F62(48), match those of the water-selective Aqp1 and AqpZ, and these residues are seen in most of the water-selective subfamily. The third corner in the selectivity filter is the main-chain

carbonyl oxygen of S196(200), which displays an electrostatic environment that is essentially equivalent to that of the carbonyl oxygen of C191(200) of Aqp1 (Fig. 3A). However, the histidine-to-isoleucine substitution results in increases in both the cross-sectional surface area and the hydrophobicity of the selectivity filter of AqpM in comparison with Aqp1 or AqpZ. At the other end of the spectrum, the aquaglyceroporin GlpF has F200 and W48 opposite R206 in its selectivity filter, which is significantly more hydrophobic than the selectivity filter of AqpM (Fig. 3D). These differences in GlpF support a selectivity filter with a large hydrophobic surface thought to aid in the passage of glycerol by providing two hydrophobic planar walls, resembling the “greasy slide” as in maltoporin (21) for the carbon–hydrogen backbone of the glycerol molecule. Therefore, the unique occurrence of an aliphatic side chain at I187(191), with its structural consequences on the size and hydrophobicity of the key selectivity filter in AqpM, represents another subfamily among the aquaporins.

The structure of AqpM reinforces the importance of the electrostatic environment and the physical size of the selectivity filter in influencing the rate of water conductance through the aquaporin channel. The aquaporin channel structures, including that of AqpM, present the paradox that an increase in the cross-sectional surface area of the selectivity filter results in a decrease in the conductance rate for water. To date, the largest selectivity filter has been seen in the relatively poor water-conducting glycerol channel GlpF. In contrast, the much more efficient water channels Aqp1 and AqpZ display the most constricted selectivity filters. These observations suggest that channel diameter and charge distribution and hydrophobicity all play a part in the mechanism of selectivity and conductance rates for aquaporins. The size of the selectivity filter of AqpM is inhibitory toward the conductance of glycerol because of steric hindrance. On the other hand, it is reasonable to conclude that the larger channel diameter of AqpM in comparison with that of Aqp1 or AqpZ results in its relatively low conductance rate of water because of the exchange of the histidine with the aliphatic isoleucine residue at the selectivity filter and the resulting loss of a hydrogen-bonding partner at this crucial location in the channel.

The Vestibules. In addition to the water and glycerol molecules coordinated in the narrow pore of the channel, a network of water and glycerol molecules on the extracellular vestibule (Fig. 4B) form a contiguous chain of hydrogen-bonded molecules that extend from the surface of the protein into the narrow portion of the channel. The glycerol molecules seen in this pocket are oriented with their greasy backbone to the hydrophobic floor of this pocket and their OH groups hydrogen-bonded to water molecules or polar atoms along the walls of the pocket. The alignment of the hydrophobic wall of the narrow channel, matched by the hydrophobic backbone of glycerol in the structure, may equally serve to recruit other hydrophobic nutrients before entering the narrow portion of the channel.

The coordination of glycerol and water molecules in the pocket of the extracellular vestibule of AqpM indicates that the selectivity and conductance of permeant molecules begins in the vestibule of the channel. The energetically unfavorable process of shedding the hydration shell of water around the permeant is initiated in the vestibule as it transitions into the channel by hydrogen-bonding interaction with waters already in the pocket. Because the hydration shell is partially stripped before the permeant enters the narrow portion of the channel, transit through the channel might thus be expedited by this role of this unique vestibule. In addition to shedding the hydration shell of the waters, the vestibule may serve to increase the local concentration of waters at the mouth of the channel, effectively increasing the gradient and therefore the rate of water conductance through the channel.

