Retinal Aquaporin-4 and Regulation of Water Inflow Into the Vitreous Body

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PURPOSE. Details of the posterior eye water dynamics are unclear. Aquaporin-4 (AQP4), a water channel, plays an important role in water dynamics in the central nervous system and is also present in the ocular tissue. The purpose of this study was to reveal the role of AQP4 in the water dynamics of the posterior eye using in vivo JJ vicinal coupling proton exchange (JJVCPE) magnetic resonance imaging (MRI) of AQP4 knockout (KO) mice and their wild-type littermates (controls).

METHODS. JJVCPE MRI of the eye was performed on five AQP4 KO mice and seven control mice. We assessed the normalized signal intensities of a region of interest (ROI) set in the vitreous body after H217O administration. The results of the two groups were compared using a two-tailed Mann-Whitney U test.

RESULTS. A statistical analysis revealed that the normalized ROI signal intensities at the steady state were significantly lower (P = 0.010, <0.05) in the AQP4 KO mice (mean ± SD, 84.5% ± 2.7%) than the controls (mean ± SD, 88.8% ± 1.9%).

CONCLUSIONS. The present study using JJVCPE MRI of the eye demonstrated that retinal AQP4 has a potential role in the regulation of water inflow into the vitreous body. Absence of AQP4 in the KO mice probably induces lower water outflow from the vitreous body. Our results could help clarify the pathogenesis of various ocular diseases.

Keywords: aquaporin-4, magnetic resonance imaging, postiridial flow, Müller cells, knockout mice

Water dynamics in the eye are still poorly understood. In its anterior part, aqueous humor exits at the anterior chamber angle through the trabecular meshwork and uveoscleral routes.1 However, water dynamics in the posterior eye have not been investigated in detail, despite indications of the existence of postiridial flow, that is, the flow of aqueous humor into the vitreous body, as previously shown by fluorescein distribution.2–4 Aquaporin-4 (AQP4) is distributed abundantly in the ocular tissue (e.g., the ciliary epithelium and Müller cells of the retina).5 Therefore, AQP4 has been suggested to be similarly involved in the water dynamics of the eye. JJ vicinal coupling proton exchange (JJVCPE) magnetic resonance imaging (MRI) has already been used to assess in vivo water dynamics in the mouse brain.5 The H217O influx, injected through the femoral vein, into the cerebrospinal fluid (CSF) can be observed as an H217O dose-dependent signal reduction in T2-weighted magnetic resonance images. In the present study, we performed JJVCPE MRI of the eyes (JJVCPE MRI of the eye) of AQP4 knockout (KO) mice and their wild-type littermates (controls) to investigate the water dynamics of the posterior eye and reveal the role of AQP4 in it.

METHODS

This study was approved and carried out in accordance with the animal research guidelines of the Institutional Review Board/Ethics Committee at Niigata University (Registration No. SA00642). This study was also performed to comply with the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research. AQP4 KO mice and controls (both sexes; 11–18 weeks old; n = 5 AQP4 KO mice; n = 7 controls) underwent JJVCPE MRI of the eye. The generation of AQP4 KO mice has been described previously.6 The mice resulted from homologous recombination using an embryonic stem cell line from the C57BL/6 strain.

JJVCPE MRI of the eyes were performed on a 15-cm bore, 7-T horizontal magnet (Magnex Scientific, Abingdon, UK) with a Varian Unity INOVA system (Varian, Palo Alto, CA, USA) equipped with an actively shielded gradient. The original JJVCPE MRI can assess in vivo water fluid dynamics in the mouse brain by setting a slab to include the CSF, the cortex, and the basal ganglia.8 For the JJVCPE MRI of the eye, a 1-mm-thick 3-dimensional region of interest (ROI) was set to include a posterior portion of the left eye, containing the vitreous body (Fig. 1A). Adiabatic double spin-echo prepared rapid acquisition with refocused echoes was utilized with the following imaging parameters: 128 × 128 matrix, 20 × 20 mm² field of view, repetition time...
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FIGURE 1. Result of a JJVCPE MRI of the eye in a control mouse and an AQP4 KO mouse. (A) A 1-mm-thick 3-dimensional (3-D) ROI was set to include a posterior portion of the left eye containing the vitreous body. The upper part shows a 3-D ROI (white rectangles) in coronal and sagittal T1-weighted images. The lower part shows the vitreous body of the left eye as donut shaped with high signal intensities in T2-weighted images by the sequences of adiabatic double spin-echo prepared rapid acquisition with refocused echoes. (B) The reduction of normalized ROI signal intensities, set to the vitreous body, after H217O administration is observed in a control mouse (upper part) and an AQP4 KO mouse (lower part) who underwent JJVCPE MRI of the eye. Normalized ROI signal intensities at steady state of a control mouse are greater than those of an AQP4 KO mouse.

2000 ms, echo train length 32, echo time (TE) for first echo 8.8 ms, echo spacing 5 ms, and effective TE 84.8 ms. The total scan time was 1 hour and 6 minutes for 500 scans per 8 seconds.

Before JJVCPE MRI of the eye, each mouse was anesthetized with inhaled isoflurane (0.01 mL/min), N2O (0.7 mL/min), and O2 (0.3 mL/min), and PE10 tubing was inserted into the right femoral vein. Using 8-0 silk, a suture was carefully made between the left bulbar conjunctiva and the left upper palpebral conjunctiva to fixate the left eye. The left upper and lower eyelids were also sutured using 8-0 silk to prevent exposure of the cornea. During JJVCPE MRI of the eye, each mouse was anesthetized with intraperitoneal administration of urethane and dexmedetomidine hydrochloride (0.6 g/kg and 0.6 g/kg, respectively) and breathed spontaneously under inhaled N2O (0.7 mL/min) and O2 (0.3 mL/min), and PE10 tubing was inserted into the right femoral vein. Using 8-0 silk, a suture was carefully made between the left bulbar conjunctiva and the left upper palpebral conjunctiva to fixate the left eye. The left upper and lower eyelids were also sutured using 8-0 silk to prevent exposure of the cornea. During JJVCPE MRI of the eye, each mouse was anesthetized with intraperitoneal administration of urethane and dexmedetomidine hydrochloride (0.6 g/kg and 0.6 g/kg, respectively) and breathed spontaneously under inhaled N2O (0.7 mL/min) and O2 (0.3 mL/min), and PE10 tubing was inserted into the right femoral vein.

The images we obtained were analyzed using image processing software (MRVision, Winchester, MA, USA). The ROI was set to the vitreous body (Fig. 1A). Normalized ROI signal intensities, expressed as percentages against the average intensity of the identical pixels before H217O administration, were plotted against time (Fig. 1B). Normalized ROI signal intensity at steady state was determined by fitting their time course by a function previously described. A two-tailed Mann-Whitney U test was used to compare the numerical data of the AQP4 KO and control mice. We selected the Mann-Whitney U test because of small sample size. Statistical analysis was done using GraphPad Prism 8 (GraphPad Software, La Jolla, CA, USA). A P value <0.05 was regarded as statistically significant.
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**RESULTS**

All 12 mice undergoing JJVCPE MRI of the eye showed a reduction in the normalized ROI signal intensity after H217O administration. The representative results of an AQP4 KO and a control mouse are shown in Figure 1B. The signal intensity was reduced, and H217O concentrations plateaued within 20 minutes.

Statistical analysis revealed that normalized ROI signal intensities at steady state were significantly lower ($P = 0.010$, $<0.05$) in the AQP4 KO mice (84.5% ± 2.7%) than in the controls (88.8% ± 1.9%) (Fig. 2).

**DISCUSSION**

In the present study, JJVCPE MRI of the eye showed that water injected through the mouse femoral vein flowed relatively rapidly into the vitreous body of the mouse eye. Due to its nuclear magnetic resonance sensitivity, 17O interferes with the neighboring 1Hs through a covalent bond in a phenomenon called JJ vicinal coupling. Accordingly, the influx of H217O into ROIs in T2-weighted magnetic resonance images causes signal reduction. Based on the results of the studies on water dynamics in the posterior eye,2–4 the water inflow into the vitreous body of the mouse eye detected in the present study probably observes a phenomenon called postiridial flow, an influx of aqueous humor into the vitreous body. In other words, H217O injected through the mouse femoral vein is thought to flow into the vitreous body as a part of the aqueous humor produced at the ciliary epithelium because no other pathways of water entry into the vitreous body have been reported aside from postiridial flow.2–4

Furthermore, we revealed that normalized ROI signal intensities at steady state were significantly lower in the AQP4 KO mice than in the controls. The results indicate that water inflow into the vitreous body was higher in the AQP4 KO mice than in the controls and/or that water outflow from the vitreous body was lower in the AQP4 KO mice than in the controls. Although the direction of the postiridial flow remains undetermined, it has been proposed that the aqueous humor produced at the ciliary epithelium passes through the vitreous body into the retina (Fig. 3).2–4 In this route, AQP4 in the ciliary epithelium and in Müller cells of the retina plays an important role. The impaired aqueous humor production at the ciliary epithelium in AQP4 KO mice induces lower water inflow into the vitreous body.10 Our results propose an additional mechanism by which Müller cells in the retina would absorb water from the vitreous body through AQP4. In AQP4 KO mice, the capacity of Müller cells of absorbing water from the vitreous body is impaired, probably leading to lower water outflow from the vitreous body (Fig. 4).

In the retina, AQP4 is mainly distributed in Müller cells and has a prominent role in fluid homeostasis in the retina.11–13 Immunohistochemical analyses have shown that AQP4 in Müller cells demonstrates colocalized distribution with Kir 4.1, which is a potassium channel.14 Kir 4.1 plays an important role in the spatial buffering of the retinal potassium, and owing to the colocalized distribution, AQP4 has a possibility of cooperating with Kir 4.1 function.15 AQP4 in Müller cells also demonstrates colocalized distribution with the transient receptor potential vanilloid 4 (TRPV4), which is a sensor for astroglial swelling.16 TRPV4 should interact with AQP4 to sense glial swelling in Müller cells and astrocytes.17

In the CNS, AQP4 is mainly distributed in the foot processes of astrocytes,18 and JJVCPE MRI has shown that water inflow into the CSF is reduced in AQP4 KO mice.8 Nakada and Kwee16 have proposed that water enters astrocytes through AQP4 and then moves into the pericapillary Virchow-Robin space. This proposal can be applied both to the water dynamics of the retina and that in the CNS. AQP4s in Müller cells are distributed in the end feet membranes.
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The glymphatic pathway between the CSF and the pericapillary space in the optic nerve using fluorescent dextran. The glymphatic pathway would be released into the pericapillary Virchow-Robin space in the retina. A recent study has shown the pathway hypothesize that water absorbed through AQP4s in Müller cells would be released into the pericapillary Virchow-Robin space facing the vitreous body and capillary endothelium. We hypothesize that water absorbed through AQP4s in Müller cells would be released into the pericapillary Virchow-Robin space in the retina. A recent study has shown the pathway between the CSF and the pericapillary space in the optic nerve using fluorescent dextran. The glymphatic pathway constituted by the perivascular flow is thought to be the clearance system of the interstitial solutes and proteins in the brain. Daruich et al. have proposed that the glymphatic pathway probably exists in the retina. We propose that the hypothesized water flow within the Müller cells, mentioned above, could drive the retinal “glymphatic pathway.”

We investigated the water dynamics in the vitreous body by administration of sufficient dose of H$_2^{17}$O (approximately 2.7 mL/kg) to detect the signal changes in the vitreous body. Among the reports of MRI studies using H$_2^{17}$O as a nonradioactive trace, Cheng et al. demonstrated the signal changes in MRI of the rabbit eye after an intravenous H$_2^{17}$O injection, wherein the signal changes were not observed in the vitreous body and rapid signal changes were observed in the anterior chamber. Although our findings are different from those reported by them, this discrepancy may be due to the difference in the dose of H$_2^{17}$O (0.1 mL/kg in the study by Cheng et al.) as they mentioned that sufficient H$_2^{17}$O should be loaded to visualize the signal changes in the vitreous body.

The present study utilizing JJVCPE MRI of the eye demonstrated that AQP4 in the retina has a potential role in the regulation of water inflow into the vitreous body via systemic circulation. Further investigation of the mechanism of aqueous humor in the posterior eye could contribute to elucidating the pathogenesis of various diseases (e.g., glaucoma and age-related macular degeneration) and the development of new treatments.

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