Vasopressin receptors V1α and V2 are not osmosensors

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Abstract
Herein, we investigated whether G protein-coupled signaling via the vasopressin receptors of the V1α and V2 subtypes (V1αR and V2R) could be obtained as a direct response to hyperosmolar challenges and/or whether hyperosmolar challenges could augment classical vasopressin-dependent V1αR signaling. The V1αR-dependent response was monitored indirectly via its effects on aquaporin 4 (AQP4) when heterologously expressed in Xenopus oocytes and V1αR and V2R function was directly monitored following heterologous expression in COS-7 cells. A tendency toward an osmotically induced, V1αR-mediated reduction in AQP4-dependent water permeability was observed, although osmotic challenges failed to mimic vasopressin-dependent V1αR-mediated internalization of AQP4. Direct monitoring of inositol phosphate (IP) production of V1αR-expressing COS-7 cells demonstrated an efficient vasopressin-dependent response that was, however, independent of hyperosmotic challenges. Similarly, the cAMP production by the V2R was unaffected by hyperosmotic challenges although, in contrast to the V1αR, the V2R displayed an ability to support alternative signaling (IP production) at higher concentration of vasopressin. V1αR and V2R respond directly to vasopressin exposure, but they do not have an ability to act as osmo- or volume sensors when exposed to an osmotic gradient in the absence or presence of vasopressin.

Introduction
The antidiuretic peptide hormone arginine vasopressin participates in the regulation of mammalian water homeostasis. It is released systemically and centrally (via vasopressinergic neurons located in direct contact with the blood vessels; Jojart et al. 1984) as a regulatory response to increases in plasma osmolarity or a reduction in blood volume (Landgraf 1992). Vasopressin exerts its effects via a family of G protein-coupled receptors. The most prominently expressed are the V1α and the V2 type (V1αR and V2R) (Lolait et al. 1995). The V1αR is found in a number of tissues including brain; it has been detected in neurons, glial cells, and endothelial cells of the blood–brain barrier (van Leeuwen et al. 1987; Chen et al. 1993; Szmydynger-Chodobska et al. 2004). The V2R has a more restricted distribution, and is predominantly expressed in the kidney (Lolait et al. 1992). Vasopressin receptors recruit different G proteins and hence confer distinct signaling pathways: activation of the V1αR leads to Gq recruitment, increased activity of phospholipase Cβ (PLCβ), generation of inositol trisphosphate (IP3), Ca2+ release from intracellular stores, and protein kinase C (PKC) activation (Zhao and Brinton 2003). Conversely, V2R-activation leads to Gs, recruitment, activation of adenylyl cyclase, and cAMP generation, followed by increased catalytic activity of cAMP-dependent protein kinases (for review, see Lolait et al. 1995). A prominent site of action for vasopressin is the kidney, in which activation of the V2R results in increased absorption of water, and ultimately helps main-
tain body water homeostasis, by modulating plasma membrane levels of the water channel AQP2 in collecting duct principal cells (Kortenoeven and Fenton 2014). Vasopressin, in addition, facilitates ion and water accumulation in the brain in a V1\(_3\)R-dependent manner (Kleindienst et al. 2006; Liu et al. 2010). We have previously demonstrated a V1\(_3\)R-mediated downregulation of aquaporin 4 (AQP4) membrane expression levels and proposed that this downregulation might be beneficial during periods of dehydration in an attempt to limit the loss of water from the brain (Moeller et al. 2009).

Since a prominent role of vasopressin receptors is to participate in volume regulation, it would be of physiological advantage for the receptor to sense the osmotic change directly in the affected cells — instead of after the appreciable delay with which vasopressin is released systemically and/or centrally. Such intrinsic osmo/volume sensing abilities have previously been described for other membrane transport proteins such as the K\(^+-\)channel KCNQ1 (Grunnet et al. 2003), the osmosensing trimeric glycine betaine transporter, BetP (Krämer and Ziegler 2009), and several ion channels of the transient receptor potential family (Plant 2014). It has thus been proposed that vasopressin-induced activation of the V1\(_3\)R could be enhanced by hyperosmolar conditions (Izumi et al. 2008). Here, we investigated whether V1\(_3\)R (or V2R)-dependent signaling could be obtained as a direct response to hyperosmolar challenges and whether hyperosmolar challenges could augment classical vasopressin-dependent V1\(_3\)R signaling.

Materials and Methods

Oocyte preparation and protein expression

*Xenopus laevis* frogs were obtained from Nasco (Fort Atkinson, WI) or National Center for Scientific Research (France). Oocytes were surgically removed from anesthetized frogs and prepared as previously described (Fenton et al. 2010). The protocol complies with the European Community guidelines for the use of experimental animals and the experiments were approved by The Danish National Committee for Animal Studies. Rat AQP4.M23 (obtained from S. Nielsen, Aalborg University, Denmark) and rat mGluR1 (obtained from J. P. Pin, Montpellier University, France) were subcloned into the oocyte expression vector pXOOM and the human V1\(_3\)R (obtained from M. J. Brownstein, NIH, Bethesda, MD) was subcloned into the vector pNB1. The cDNAs were linearized downstream from the poly-A segment and in vitro transcribed using T7 mRNA Machine (Ambion, Austin, TX). The cRNA was then extracted with MEGAclear (Ambion, Austin, TX) and microinjected into defolliculated *Xenopus* oocytes (8 ng rAQP4 RNA/oocyte, 16 ng hV1\(_3\)R RNA/oocyte, or 16 ng rmGluR1a RNA/oocyte). The oocytes were kept in Kulori medium (in mmol/L: 90 NaCl, 1 KCl, 1 CaCl\(_2\), 1 MgCl\(_2\), 5 HEPES, pH 7.4) for 4–6 days at 19°C prior to experiments.

Volume measurements

The experimental setup for measuring water permeability of oocytes has been described in detail previously (Zeuthen et al. 2006). Briefly, the oocyte was placed in a small chamber with a glass bottom and perfused with a control solution (in mmol/L: 100 NaCl, 2 KCl, 1 CaCl\(_2\), 1 MgCl\(_2\), 10 HEPES, pH 7.4) at room temperature. Images of the oocytes were captured continuously from below at a rate of 25 images/sec. The oocytes were challenged with a hyperosmolar solution (control solution containing an additional 50 mOsm mannitol) in order to determine the osmotic water permeability:

\[
L_p = \frac{-J_w}{A \times \Delta \pi \times V_W}
\]

where \(J_w\) is the initial water flux during osmotic challenge, \(A\) is membrane surface area (nine times the apparent surface area due to membrane folding: Zampighi et al. 1995), \(\Delta \pi\) is the osmotic challenge, and \(V_W\) is the partial molal volume of water (18 cm\(^3\)/mol). All osmolarities were validated using a cryoscopic osmometer Type 15 (Löser, Berlin, Germany).

Immunocytochemistry

Oocytes (3–6 per condition per experiment) were fixed for 1 h in 3% paraformaldehyde in Kulori medium, rinsed in Kulori medium, dehydrated in a series of ethanol concentrations (30 min in 70%, 96%, and 99%) followed by incubation in xylene for 1 h. Oocytes were infiltrated with paraffin for 1 h at 50°C before embedding. Sections of 2 μm were cut on a Leica RM 2126 microtome and immunostaining performed as described previously (Fenton et al. 2010) using a rabbit polyclonal anti-AQP4 antibody 1:5000 (Alamone Laboratories, Jerusalem, Israel). An Alexa 488-conjugated secondary antibody 1:1,000 was used for visualization (DAR; Invitrogen, Nærum, Denmark). A Leica TCS SL confocal microscope and Leica confocal software were used for imaging of the oocytes. Control AQP4-expressing oocytes were used to set laser intensity and capture settings on the microscope such that saturation of images for each condition was avoided. The microscope and laser settings were kept constant for imaging of each oocyte within each experiment. Images were taken using an HCX PL APO ×63 oil objective lens (numerical aperture: 1.40). A minimum of two
images per oocyte, with 3–6 oocytes per experiment, were used for statistical analysis. Image semiquantification and validation was performed as previously described (Fenton et al. 2010).

Cell culture and transfection
COS-7 cells were grown in Dulbecco’s modified Eagle’s medium (Invitrogen) supplemented with 10% fetal calf serum, 2 mmol/L glutamine, 180 U/mL penicillin, and 45 μg/mL streptomycin (10% CO2, 37°C). Receptor signaling experiments were carried out using COS-7 cells transfected with the human vasopressin receptors, V1aR or V2R in pcDNA3 (obtained from M. J. Brownstein, NIMH, Bethesda, MD). The transfection was performed using the calcium phosphate precipitation method (Graham and Van der Eb 1973) according to Kissow et al. (2012).

Inositol phosphate assay
Inositol phosphate accumulation was measured using a scintillation proximity assay (SPA). One day after transfection, COS-7 cells were seeded at 35,000 cells/well in 96-well plates for 24 h with 0.5 μCi of myo-[3H]inositol (NET1156, PerkinElmer, Waltham, MA) in 100 μL medium per well. Cells were washed twice in Hank’s balanced salt solution (HBSS, in mmol/L: 138 NaCl, 4 NaHCO3, 0.33 Na2HPO4, 5 KCl, 0.4 KH2PO4, 1.3 CaCl2, 0.5 MgCl2, 0.4 MgSO4, and 5.5 D-glucose) prior to 15 min incubation in HBSS supplemented with 10 mmol/L LiCl at 37°C to block IP1 breakdown. Subsequently cells were treated with vasopressin (in concentrations as indicated in the text) or hyperosmolar challenges (in duplicate or triplicate determinations) for 90 min at 37°C. Cells were treated with the enzyme fragment complementation-based cAMP assay according to manufacturer’s instructions (HitHunter cAMP XS assay, DiscoveRx, Fremont, CA). The cAMP content was measured as luminescence using an EnVision 2104 Multitable Platereadper (PerkinElmer) with a cAMP standard curve for validation.

Statistics
EC50 values were obtained from dose–response curves obtained with fitting to log(agonist) versus response with variable slope (four parameters) according to the equation

\[
Y = \frac{\text{Bottom} + (\text{Top} - \text{Bottom})}{1 + 10^{(\text{LogEC}_{50} - X) \times \text{HillSlope})}}
\]

in GraphPad Prism 6.0 software (San Diego, CA). Statistical analyses were carried out in GraphPad Prism 6.0 software as described in the text. Data were obtained from at least three different animal donors (Xenopus laevis) or at least three different cell transfections and are presented as means ± SEM (confidence intervals [CI] when Log scales were employed). A probability level of \( P < 0.05 \) was considered statistically significant.

Results
Putative osmosensing by V1aR assessed by water permeability measurements
Activation of the Gq protein-coupled vasopressin receptor V1aR expressed in Xenopus laevis oocytes initiates a signaling cascade involving phospholipase C and increased levels of intracellular Ca2+ (Nathanson et al. 1992; Ancelin and Morel 1998). We have previously in the Xenopus laevis expression system, by several complementary technical approaches, demonstrated that activation of V1aR led to an internalization of coexpressed AQP4 and thus a reduction in the osmotic water permeability of the oocytes (Moeller et al. 2009). We therefore initially utilized this experimental setup to obtain a functional readout for V1aR activation with the aim to determine if cell
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Figure 1. V1aR-dependent downregulation of AQP4. (A) Volume traces obtained from an uninjected oocyte (left panel) and an AQP4/V1aR-expressing oocyte challenged with an osmotic gradient of 50 mOsm mannitol for 30 sec. (B) Relative water permeability of AQP4/V1aR-expressing oocytes (n = 20) was reduced significantly more than those expressing only AQP4 (n = 5) after exposure to vasopressin for 30 min (P < 0.05, Fig. 1B), indicating that activation of the receptor leads to downregulation of AQP4-mediated water permeability. To test if V1aR could directly respond to hyperosmotic challenges, AQP4/V1aR-expressing oocytes (n = 22) and AQP4-expressing oocytes (n = 6) were exposed to consecutive hyperosmotic challenges, as above, but in the absence of vasopressin in the test medium (Fig. 1C). The osmotic water permeability of the oocytes expressing either AQP4/V1aR or AQP4 did not differ in their response to repeated hyperosmotic challenges, although a tendency toward a difference between the two groups was observed (P = 0.053). The low inherent water permeability of the uninjected oocyte was not significantly affected by exposure to vasopressin or to repeated osmotic challenges (data not shown). Due to the observed tendency for an osmotically induced reduction in the water permeability of AQP4/V1aR-expressing oocytes, a parallel control experimental series was performed with an unrelated Go protein-coupled glutamate receptor (mGluR1) to determine if Go protein-coupled receptors shared a general feature of hyperosmolarity-induced Go protein activation. Similar to that observed with V1aR, the water permeability of AQP4/mGluR1-expressing oocytes was reduced upon activation of mGluR1, by addition of glutamate (500 μmol/L) to the test medium, to a level significantly different from exposing the AQP4/mGluR1 to repeated hyperosmolar challenges in the absence of glutamate (n = 8, P < 0.05, Fig. 1D). These data indicate that agonist-induced activation of Go-

shrinkage in itself can mimic previously published vasopressin-dependent activation of V1aR in the oocytes (Moeller et al. 2009). AQP4 was expressed in the oocytes either alone (AQP4) or along with V1aR (AQP4/V1aR). The oocytes were monitored for volume changes with a sensitive camera while abruptly challenged with a hyperosmotic gradient of 50 mOsm (obtained by addition of 50 mmol/L mannitol to the test solution to increase the osmolarity while keeping the ionic strength constant). The rate of osmotically induced oocyte shrinkage increased dramatically upon expression of AQP4 (~15-fold over that of uninjected control oocytes) and amounted to a physiologically relevant cell shrinkage of around 1% during exposure to the osmotic challenge, see representative volume traces in Figure 1A. After three initial control water permeability measurements, the oocytes were exposed to saturating concentrations of vasopressin to obtain maximal activation of V1aR (1 μmol/L, indicated by the black bar in Fig. 1B) and the osmotic water permeability determined as a function of time and normalized to the control value (Fig. 1B). The water permeability of AQP4/V1aR-expressing oocytes (n = 20) was reduced significantly more than those expressing only AQP4 (n = 5) after exposure to vasopressin for 30 min (P < 0.05, Fig. 1B), indicating that activation of the receptor leads to downregulation of AQP4-mediated water permeability. To test if V1aR could directly respond to hyperosmotic challenges, AQP4/V1aR-expressing oocytes (n = 22) and AQP4-expressing oocytes (n = 6) were exposed to consecutive hyperosmotic challenges, as above, but in the absence of vasopressin in the test medium (Fig. 1C). The osmotic water permeability of the oocytes expressing either AQP4/V1aR or AQP4 did not differ in their response to repeated hyperosmotic challenges, although a tendency toward a difference between the two groups was observed (P = 0.053). The low inherent water permeability of the uninjected oocyte was not significantly affected by exposure to vasopressin or to repeated osmotic challenges (data not shown). Due to the observed tendency for an osmotically induced reduction in the water permeability of AQP4/V1aR-expressing oocytes, a parallel control experimental series was performed with an unrelated Go protein-coupled glutamate receptor (mGluR1) to determine if Go protein-coupled receptors shared a general feature of hyperosmolarity-induced Go protein activation. Similar to that observed with V1aR, the water permeability of AQP4/mGluR1-expressing oocytes was reduced upon activation of mGluR1, by addition of glutamate (500 μmol/L) to the test medium, to a level significantly different from exposing the AQP4/mGluR1 to repeated hyperosmolar challenges in the absence of glutamate (n = 8, P < 0.05, Fig. 1D). These data indicate that agonist-induced activation of Go-
coupled receptors confers downregulation of AQP4-mediated water permeability which was, however, not significantly mimicked by exposure of the oocytes to repeated hyperosmolar challenges.

**Putative osmosensing by V1\textsubscript{a}R assessed by immunocytochemistry**

To assess a role of hyperosmotic challenge on the V1\textsubscript{a}R using an alternative approach, we analyzed the membrane abundance of AQP4 in oocytes. We have previously shown a V1\textsubscript{a}R-dependent internalization of AQP4 following vasopressin exposure (Moeller et al. 2009) and aimed in the present experimental series to determine if the vasopressin-dependent internalization of AQP4 could be mimicked by cell shrinkage. AQP4/V1\textsubscript{a}R-expressing oocytes were exposed to either vasopressin or repeated hyperosmotic challenges and the plasma membrane abundance of AQP4 subsequently quantified by immunocytochemistry. AQP4/V1\textsubscript{a}R-expressing oocytes were exposed to either vasopressin or repeated hyperosmotic challenges and the plasma membrane abundance of AQP4 subsequently quantified by immunocytochemistry. Oocytes expressing AQP4 alone or AQP4/V1\textsubscript{a}R were incubated for 80 min in control conditions, in the presence of 1 \textmu mol/L vasopressin for the last 60 min, or challenged with an extracellular osmotic gradient of 50 mOsm mannitol for 30 sec every 10 min during the span of the 80 min incubation time (designed to mimic the water permeability measurements from Fig. 1).

Representative confocal images of AQP4- and AQP4/V1\textsubscript{a}R-expressing oocytes are shown in Figure 2A. No immune reactivity toward AQP4 was detected in uninjected oocytes (data not shown and Moeller et al. 2009). The plasma membrane abundance of AQP4 was quantified and normalized to the membrane abundance observed under control conditions in AQP4- and AQP4/V1\textsubscript{a}R-expressing oocytes, respectively (Fig. 2B). Exposure to vasopressin induced a reduction in plasma membrane abundance in AQP4/V1\textsubscript{a}R-expressing oocytes (55 ± 12% of control, \(n = 5\) experiments with 3–6 oocytes per condition) which was significantly more than that obtained in AQP4-expressing oocytes (95 ± 11% of control, \(n = 5\) experiments with 3–6 oocytes per condition), \(P < 0.05\). The vasopressin-induced AQP4 internalization in the AQP4/V1\textsubscript{a}R-expressing oocytes was not mimicked by repeated hyperosmolar challenges (compare 92 ± 5% of control for hyperosmolar challenges and the 55 ± 12% of control for vaso-

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**Figure 2.** V1\textsubscript{a}R-dependent internalization of AQP4. (A) Confocal laser scanning microscopy of oocytes expressing either AQP4 (left panel) or AQP4/V1\textsubscript{a}R (right panel) immune-labeled for AQP4. The upper panels are representative images of oocytes exposed to control solution without vasopressin for 80 min. The middle panels are representative images of oocytes kept in control solution for 20 min and then treated with 1 \(\mu\)mol/L vasopressin for 60 min. The lower panels are representative images of oocytes treated with a 50 mOsm hyperosmolar gradient for 30 sec every 10 min of an 80-min incubation period. (B) Oocyte plasma membrane fluorescence intensity normalized to that of the oocytes kept in control solution, \(n = 5\) experiments with 3–6 oocytes per condition. The indicated groups were compared using one-way analysis of variance (ANOVA) with Sidák’s multiple comparison post hoc test. *\(P < 0.05\); ns, not significant.
pressin, \( n = 5 \) experiments with 3–6 oocytes per condition, \( P < 0.05 \). Oocytes expressing AQP4 alone, likewise, did not respond to repeated hyperosmolar challenges (104 ± 6\% of control, \( n = 5 \) experiments with 3–6 oocytes per condition). In the present experimental setting, based on an indirect experimental read-out of \( V_{1a}R \) function, the \( V_{1a}R \) responds to vasopressin exposure but not to hyperosmotic challenges, indicating that \( V_{1a}R \) does not act as an osmosensor.

**Putative osmosensing by \( V_{1a}R \) assessed by quantification of intracellular secondary messengers**

Activation of \( V_{1a}R \) by vasopressin leads to classical \( G_{q} \) protein recruitment and induction of a downstream intracellular signaling cascade involving phospholipase C activation, IP\(_3\) production, \( Ca^{2+} \) release from intracellular stores and associated PKC activation (Thibonnier et al. 1994). To obtain a direct experimental read-out of \( V_{1a}R \) activity in mammalian cells, COS-7 cells were transiently transfected with the \( V_{1a}R \) and \( V_{1a}R \)-mediated IP accumulation upon exposure to vasopressin and/or hyperosmotic challenges was assessed. Functional expression of the \( V_{1a}R \) was initially verified by performing a vasopressin dose–response curve of \( V_{1a}R \)-transfected cells versus vector-transfected control cells. Application of vasopressin had no effect on IP production in vector-transfected control cells but significantly increased the IP accumulation in a dose-dependent manner (EC\(_{50} = 0.68 \text{ nmol/L, 95\% CI: } 0.44; 1.05, n = 7, \text{ Fig. 3A}) to 9.4 ± 1.9-fold in \( V_{1a}R \)-transfected cells, significantly higher than that observed in vector-transfected cells

**Figure 3.** Hyperosmolar effects on \( V_{1a}R \)-dependent inositol phosphate (IP) production. (A) Vasopressin dose–response curve measured with IP production in COS-7 cells transfected with either vector alone or with \( V_{1a}R, n = 7 \). (B) IP production of COS-7 cells transfected with either vector alone or with \( V_{1a}R \) upon addition of vasopressin (1 \( \mu \text{mol/L} \)) or various hyperosmolar treatments as indicated on graph, \( n = 3–8 \). (C) Effect of hyperosmolar treatment (50 or 150 mOsm) on the vasopressin dose–response curve, \( n = 3–8 \). (D) cAMP production obtained in COS-7 cells transfected with either vector alone or with \( V_{1a}R \) in response to vasopressin exposure (1 \( \mu \text{mol/L} \)) or various hyperosmolar treatments as indicated on graph, \( n = 4–6 \). Data obtained with \( V_{1a}R \)-transfected cells were compared to those obtained with vector-transfected cells using two-way analysis of variance (ANOVA) with Sidák’s multiple comparison post hoc test and indicated with * (note: the difference between the vasopressin-induced cAMP production in vector- and \( V_{1a}R \)-transfected cells was not statistically significant even when compared with Student’s t-test). Statistical significance of vasopressin-induced cAMP production in vector-transfected cells was determined with two-way ANOVA with a Dunnett’s multiple comparison test and indicated with #. *\( P < 0.05; ***P < 0.001; \text{ ns, not significant.}

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the vector-transfected cells, \( n = 3–8 \), \( P < 0.001 \) (Fig. 3B). To determine a putative hyperosmolar-induced activation of V1\(_1\)R, vector- or V1\(_1\)R-transfected cells were, in parallel, exposed to a 50 mOsm hyperosmotic challenge either (1) for 30 sec followed by return to control solution, (2) for the entire accumulation period (90 min), or (3) as a gradual 5 mOsm increase every 10 min (Fig. 3B). None of these hyperosmolar challenges promoted significant V1\(_1\)R activation as measured by IP accumulation (\( n = 3–8 \) experiments), indicating a lack of receptor-dependent osmosensing.

It was previously suggested that hyperosmolarity could augment the V1\(_1\)R response to vasopressin (Izumi et al. 2008). To assess a putative synergistic effect of vasopressin and hyperosmolar challenges on the V1\(_1\)R, dose–response curves for IP production were generated in V1\(_1\)R-transfected cells following vasopressin exposure in isosmolar assay solutions or in solutions containing hyperosmolar loads of 50 or 150 mOsm in the assay solution (Fig. 3C). 50 mOsm hyperosmotic assay solutions lowered the \( E_{\text{max}} \) (measured at 1 nmol/L AVP) to 93 ± 2% (\( n = 4 \), \( P < 0.05 \)), whereas addition of 150 mOsm lowered \( E_{\text{max}} \) to 76 ± 5% (\( n = 3 \), \( P < 0.001 \)) compared to isosmotic vasopressin exposure (assessed with one-way analysis of variance [ANOVA] with Dunnett’s multiple comparison post hoc test). Therefore, rather than a hyperosmolar-induced increase in V1\(_1\)R activity, the combined effect of vasopressin exposure in a hyperosmolar setting resulted in decreased V1\(_1\)R activity. Hyperosmotic challenges did, in addition, significantly increase the potency of vasopressin on the V1\(_1\)R: compare EC\(_{50}\) of 0.31 nmol/L, 95% CI: 0.22; 0.43, \( n = 4 \) obtained in the presence of 50 mOsm hyperosmotic challenge with the EC\(_{50}\) of 0.68 nmol/L, 95% CI: 0.44; 1.05, \( n = 7 \) obtained in isosmotic control solution, \( P < 0.05 \).

A range of G protein-coupled receptors display biased signaling, that is, they are capable of signaling through various G proteins and/or other intracellular signaling cascades depending on the specific agonist (Violin and Lefkowitz 2007; Steen et al. 2014). To determine if the V1\(_1\)R, upon a hyperosmolar stimulus, was capable of signaling through an alternative canonical signaling pathway involving G\(_i\) recruitment, activation of adenylate cyclase and increased cAMP production, vector- and V1\(_1\)R-transfected cells were exposed to vasopressin or to hyperosmolar challenge and intracellular cAMP levels measured. The V1\(_1\)R did not promote cAMP production under any of the experimental conditions (\( n = 4–6 \) experiments, Fig. 3D), suggesting that the V1\(_1\)R is unable to couple to G\(_i\) following vasopressin exposure or hyperosmolar challenges. We did, however, note a slight vasopressin-induced increase in cAMP production in vector-transfected cells (\( n = 5 \), \( P < 0.05 \)), indicating the presence of endogenous V2R in the COS-7 cells. This endogenous cAMP production was, however, not sensitive to hyperosmolarity either and did therefore not interfere with our interpretation. These results indicate that the V1\(_1\)R is unable to couple to G\(_i\) under the present experimental conditions and that it reacts solely to vasopressin exposure, resulting in G\(_i\) activation, in a manner only slightly affected by simultaneous hyperosmolar challenges in the physiological range.

**Putative osmosensing by V2R assessed by quantification of intracellular secondary messengers**

The V2R is related to the V1\(_1\)R, but is abundantly expressed in the kidney where it signals predominantly via activation of G\(_i\), resulting in increased adenylate cyclase activity and increased cAMP production (Juul et al. 2014). Due to its expression in the kidney collecting duct, the V2R faces a progressively increasing hyperosmotic environment in the transition from cortical to medullary segments, suggesting that hyperosmolarity may regulate V2R function. Thus, in a separate experimental series, the response of the V2R to direct hyperosmotic challenge was determined. COS-7 cells were transiently transfected with the V2R or empty vector and vasopressin-dependent cAMP production was determined. A slight cAMP production was observed in vector-transfected cells upon addition of vasopressin to the assay solution (\( P < 0.05 \), \( n = 5 \)), although in V2R-transfected cells, the cAMP production increased in a dose-dependent manner (Fig. 4A) to 4.6 ± 0.9-fold, which is significantly higher than that observed in the vector-transfected cells, \( n = 5 \), \( P < 0.001 \), Fig. 4B. The V2R displayed higher potency toward vasopressin (EC\(_{50}\) = 0.068 nmol/L, 95% CI: 0.044; 0.107, \( n = 5 \)) than the V1\(_1\)R, \( P < 0.001 \). To determine a putative hyperosmolar-induced activation of the V2R, vector- or V2R-transfected cells were, in parallel, exposed to a 50 mOsm hyperosmotic challenge either (1) for 30 sec followed by return to control solution, (2) for the entire accumulation period (30 min), or (3) as a gradual 10 mOsm increase every 6 min (Fig. 4B). None of these hyperosmolar challenges promoted V2R activation as measured by cAMP production (\( n = 4–6 \) experiments), indicating lack of receptor-dependent osmosensing.

To determine whether the V2R was capable of signaling through the alternative G\(_i\) signaling pathway (i.e., the main signaling pathway for the V1\(_1\)R), IP accumulation was determined in COS-7 cells transiently expressing the V2R. At vasopressin concentrations above 1 nmol/L, IP production of the V2R-expressing COS-cells was increased (black symbols in Fig. 4C, included for comparison is a dashed line for cAMP production from Fig. 4A). The maximal IP
production obtained with the V2R-expressing COS-7 cells amounted to 33 ± 3% of that obtained with V1aR-expressing COS-7 cells, n = 3, P < 0.01 (assessed with Student’s t-test) and the potency toward vasopressin was significantly reduced (compare EC₉₀ = 9.5 nmol/L, 95% CI: 6.5; 13.7, n = 7 for IP production with EC₉₀ = 0.068 nmol/L, 95% CI: 0.044; 0.107 for cAMP production, n = 5, P < 0.001). Simultaneous hyperosmolar hyperbaric challenge of 50 mOsm did not affect the IP production of the vasopressin dose–response relationship (gray symbols in Fig. 4C): compare EC₉₀ of 5.5 nmol/L, 95% CI: 3.8; 8.1, n = 4 obtained in the presence of 50 mOsm hyperosmolar challenge with the EC₉₀ of 9.5 nmol/L, 95% CI: 6.5; 13.7, n = 7 obtained in isosmotic control solution, P = 0.20.

**Discussion**

An ability of specific vasopressin receptors to respond to local hyperosmotic stress, rather than exclusively being activated by increases in systemically circulating vasopressin would theoretically placate the delay inherent in hyperosmotic-induced vasopressin release. Such “osmosensors” have been detected both peripherally and centrally, mainly in the organum vasculosum of lamina terminalis (OVLT) (Ciura and Bourque 2006; Bourque 2008). The neurons of OVLT respond readily to dehydration through cellular shrinkage and mechanically induced increase in channel opening of the transient receptor potential vanilloid 1 (TRPV1) channel, which via neuronal depolarisation leads to systemic and/or central release of vasopressin (Ciura and Bourque 2006; Ciura et al. 2011). Since a prominent role of the vasopressin receptors is to orchestrate cellular and systemic volume regulation, it would be of physiological advantage for the receptors to, in addition, sense the osmotic change directly in the affected cells and initiate the cellular response on the onset of systemic dehydration. In fact, a previous study of V1aR-expressing cells reported a marginal release of intracellular Ca²⁺ in response to a hyperosmotic challenge in the absence of vasopressin and a substantial increase in vasopressin-induced intracellular Ca²⁺ release upon vasopressin exposure during a hyperosmotic challenge (Izumi et al. 2008). Here we have resolved whether vasopressin receptors have the ability to respond directly to hyperosmotic challenge by examining V1aR-dependent effects upon heterologous expression in Xenopus oocytes or in mammalian cells. These cell types were chosen to obtain a cellular system in which the activity of the vasopressin receptors could be determined in isolation with limited contribution of other membrane channels, transporters, or receptors which could potentially display unidentified osmosensing abilities. We aimed to obtain cell shrinkage at a physiological relevant, yet experimentally detectable, level, given the relatively limited osmotic water permeability of Xenopus oocytes and COS7 cells, compared to that of the osmosensing brain areas. We therefore opted for a hyperosmotic challenge of 50 mOsm which we demonstrated to provide around 1% cell shrinkage of the oocytes and predicted to provide a maximum of 15% shrinkage of the COS7 cells. By examining V1aR-dependent effects on AQP4 function or by quantifying V1aR activity both directly (IP production) and indirectly (downstream protein regulation), we observed an absolute requirement for vasopressin to activate the V1aR. However, neither osmotic induction of receptor activity nor an osmotic augmentation of the

**Figure 4.** Hyperosmotic effects on V2R-dependent cAMP and inositol phosphate (IP) production. (A) Vasopressin dose–response curve measured with cAMP response of COS-7 cells transfected with either vector alone or with V2R, n = 5. (B) cAMP production of COS-7 cells transfected with either vector alone or with V2R upon addition of vasopressin (1 μmol/L) or various hyperosmolar treatments as indicated on graph, n = 4–6. (C) IP production (black and gray circles) and cAMP accumulation (dashed line, adapted from panel A) of COS-7 cells transfected with V2R upon vasopressin exposure, n = 4–7. Data obtained with V2R-transfected cells were compared to those obtained with vector-transfected cells using two-way ANOVA with Šidák’s multiple comparison post hoc test and indicated with * Statistical significance of vasopressin-induced cAMP production in vector-transfected cells was determined with two-way ANOVA with a Dunnett’s multiple comparison test and indicated with #. *P < 0.05; **P < 0.01; ns, not significant.
vasopressin-dependent response (at any vasopressin concentration) was observed, although the potency toward vasopressin was slightly increased during an osmotic challenge. These findings contrast those of Izumi and colleagues who employed intracellular Ca\textsuperscript{2+} measurements as their functional read-out of a putative synergistic effect of vasopressin and hyperosmolality on V1\textsubscript{a}R (Izumi et al. 2008). Although we have not identified the experimental reason for the discrepancy with this study, it is possible that the excessively large hyperosmolar gradient (\(\Delta 200\) mOsm) employed in the previous study promoted cellular signaling which would not be apparent at more physiological levels of cell shrinkage. Alternatively, one may speculate whether the open probability of putative endogenously expressed volume-sensitive Ca\textsuperscript{2+} channels, such as TRP-channels, could be modulated by activation of V1\textsubscript{a}R and thereby indirectly generate the observed synergistic effect of hyperosmolarity and vasopressin on the Ca\textsuperscript{2+} signaling. In favor of alternative direct mechanisms of osmosensing, hyperosmolar perfusion directly increased the water permeability of rat inner medullary collecting duct (Nadler 1990; Kudo et al. 1992) even in the absence of vasopressin (Kudo et al. 1992).

V1\textsubscript{a}R activation classically results in G\textsubscript{s} protein recruitment, PLC\textbeta\textsubscript{2} activation, inositol triphosphate (IP\textsubscript{3}) generation, and subsequent release of Ca\textsuperscript{2+} from intracellular stores (Thibonnier et al. 1994). However, V1\textsubscript{a}R-dependent excitation of motoneurons in a Ca\textsuperscript{2+}-, PLC\textbeta\textsubscript{2}-, and PKC-independent manner has been demonstrated (Reymond-Marron et al. 2006) and the observed V1\textsubscript{a}R-dependent cAMP production was proposed to originate from G\textalpha signaling (Wrobel et al. 2011). However, in the present study neither exposure of the V1\textsubscript{a}R to vasopressin nor hyperosmotic stress resulted in alternative signaling through the adenylate cyclase signaling pathway. Although these data indicate that the V1\textsubscript{a}R cannot signal through G\textalpha in our experimental system, we cannot exclude that V1\textsubscript{a}R-dependent activation of phospholipase A\textsubscript{2} and phospholipase D, or direct signaling via the beta-arrestin pathway may act as alternative signal transduction pathways for the V1\textsubscript{a}R (Briley et al. 1994; Terrillon et al. 2004).

Similar to the V1\textsubscript{a}R, the V2R was dependent on the presence of vasopressin for receptor activation and was insensitive to osmotic challenges in the absence of vasopressin. We confirmed the previously reported ability of the V2R to activate G\textsubscript{q} (Zhu et al. 1994) although with a lower IP production than that of V1\textsubscript{a}R, which may be due to either lower V2R-mediated coupling efficiency toward G\textsubscript{q} or to lower expression levels of V2R, both compared to that of V1\textsubscript{a}R. V2R displayed lower potency toward vasopressin when signaling through the G\textsubscript{q} protein than following its classical interaction with G\textalpha (140-fold (this study); 35-fold (Zhu et al. 1994)). The V2R-dependent signaling via G\textsubscript{q} was, as was the case for G\textalpha signaling, unperturbed by a simultaneous hyperosmolar challenge. V2R-mediated G\textsubscript{q}-signaling thus occurs exclusively at excessively high vasopressin concentrations.

In conclusion, the V1\textsubscript{a}R and the V2R respond directly to vasopressin exposure and do not share an ability to act as osmo- or volume sensors when exposed to an osmotic gradient in the absence or presence of vasopressin. At higher vasopressin concentrations, the V2R displayed G\textsubscript{q}-dependent signaling, whereas the V1\textsubscript{a}R failed to support G\textalpha-dependent signaling at the tested ligand concentrations. Neither the V1\textsubscript{a}R nor V2R induced any signaling in response to cell shrinkage. Taken together, vasopressin receptor-mediated cell volume regulation is not initiated via cell shrinkage but rather taking place exclusively upon systemic and/or central release of vasopressin during dehydration.

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**Conflicts of Interest**

None declared.

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