Investigation of LRRC8-Mediated Volume-Regulated Anion Currents in Xenopus Oocytes

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ABSTRACT Volume-regulated anion channels (VRACs) play an important role in controlling cell volume by opening upon cell swelling. Recent work has shown that heteromers of LRRC8A with other LRRC8 members (B, C, D, and E) form the VRAC. Here, we used Xenopus oocytes as a simple system to study LRRC8 proteins. We discovered that adding fluorescent proteins to the C-terminus resulted in constitutive anion channel activity. Using these constructs, we reproduced previous findings indicating that LRRC8 heteromers mediate anion and osmolyte flux with subunit-dependent kinetics and selectivity. Additionally, we found that LRRC8 heteromers mediate glutamate and ATP flux and that the inhibitor carbenoxolone acts from the extracellular side, binding to probably more than one site. Our results also suggest that the stoichiometry of LRRC8 heteromers is variable, with a number of subunits \( R \geq 6 \), and that the heteromer composition depends on the relative expression of different subunits. The system described here enables easy structure-function analysis of LRRC8 proteins.

INTRODUCTION

As a consequence of volume changes, cells activate regulatory mechanisms involving several transporters and ion channels to restore their original size (1). These mechanisms are named regulatory volume increase and regulatory volume decrease (RVD) (2). The process of RVD relies on a swelling-induced increase in the activity of a volume-regulated anion channel (VRAC) that releases organic osmolytes and anions, accompanied by the release of \( K^+ \) ions through potassium channels, and followed by osmotically driven water efflux. The anion channel has also been called the volume-sensitive organic osmolyte-anion channel (VSOAC) or the volume-sensitive outwardly rectifying anion channel (VSOR) (3). We will refer to it in the following as the VRAC.

The biophysical properties of VRACs have been extensively studied (3–6). VRAC-mediated currents activate slowly under extracellular hypotonic challenge showing a moderate instantaneous outward rectification, which reflects voltage dependence of the single-channel conductance and inactivation at positive voltages (7). Nonspecific inhibitors, such as DCPiB (8,9) or carbenoxolone (CBX) (10), which is also a gap junction blocker (11,12) and pannexin blocker (13), are found to block VRACs.

The molecular identification of the proteins responsible for VRAC activity has been finally achieved after a long history of false positives (14,15). Two groups independently identified heteromers of the leucine-rich-repeat-containing 8, member A (LRRC8A) protein with other members of the LRRC8 family (B, C, D, and E) as components of VRACs (16,17). LRRC8B–E require coexpression with LRRC8A for plasma membrane delivery (17). The subunit composition of LRRC8 heteromers affects some channel properties, such as depolarization-dependent inactivation (18), substrate specificity (17,19), and single-channel conductance (20).

Investigating the functional properties of VRAC activity induced by LRRC8 proteins is challenging. VRAC currents have been found in most vertebrate cell lines examined (14). To study LRRC8A, it was necessary to complement siRNA-depleted (16) or genome-edited cell lines (17), as the
overexpression of LRRCA suppressed the endogenous VRAC activity in cell lines by an unknown mechanism (16,17). Furthermore, the study of LRRC8 heteromers required knocking out all five LRRC8 genes by genome editing (17,20). More recently, LRRC8 heteromers have also been studied after functional reconstitution in droplet lipid bilayers (20). In addition, LRRC8 currents activate slowly by hypotonicity (17), resulting in nonsteady current levels, which complicates quantitative analysis.

In this work, we describe an alternative approach to functionally characterize VRAC activity induced by LRRC8 proteins in *Xenopus* oocytes. With this approach, we were able to perform a detailed biophysical characterization including subunit dependence of current kinetics and ion selectivity, single-channel conductance, osmolyte influx, ATP efflux, inhibitor sensitivity, osmolarity dependence, and stoichiometry.

**MATERIALS AND METHODS**

**Molecular biology**

Plasmids presented herein were constructed using standard molecular biology techniques employing recombinant polymerase chain reaction and the Multisite Gateway system (Invitrogen, Carlsbad, CA). The integrity of all constructs was confirmed by DNA sequencing.

For expression in *Xenopus* oocytes, after linearization by NotI, complementary RNA (cRNA) of human LRRC8 proteins was transcribed using the mMessage mMachine SP6 kit (Ambion, Waltham, MA). We produced cRNA of LRRCA/B/C/D/E constructs and hLRRC8A/E-VFP and hLRRC8A/B/C/D/E-monomeric Cherry (mCherry) fluorescent constructs. They were expressed by themselves or coexpressed mainly with LRRCA or LRRCA-VFP.

**Oocyte preparation and injection**

Oocytes were harvested from *Xenopus laevis* frogs that had been anesthetized by tricaine (ethyl 3-aminobenzoate methanesulfonate salt; Sigma Aldrich, St. Louis, MO) at a concentration of 1.5 gL⁻¹ buffered to neutral pH with sodium bicarbonate. After surgery, frogs were allowed to recover from anesthesia, and suitable aftercare was given. All animal protocols conformed to the European Community Guidelines on Animal Care and Experimentation and were approved by the Ethics Committee for Animal Experimentation of the University of Barcelona for the experiments conducted in Genoa. Oocytes were enzymatically defolliculated by an ~1 h treatment with collagenase type I A from *Clostridium histolyticum* (Sigma Aldrich) in a solution containing (in mM) 90 NaCl, 2 KCl, 1 MgCl₂, and 10 Heps at pH 7.5 with gentle shaking at room temperature. Fifty nanoliters of cRNA, normally containing 12 ng of each subunit, was injected with a microinjector (Nanoject II, Drummond Scientific, Broomall, PA).

**Electrophysiology (voltage clamp)**

After cRNA injection, oocytes were maintained at 18°C in a solution containing (in mM) 90 NaCl, 2 KCl, 1 MgCl₂, 1 CaCl₂, and 10 Heps at pH 7.5 or in modified Barth’s solution that contained (in mM) 88 NaCl, 1 KCl, 0.41 CaCl₂, 0.82 MgSO₄, 0.33 Ca(NO₃)₂, 2.4 NaHCO₃, and 10 Heps at pH 7.4 containing 10 mg/mL of gentamicin. One to three days after injection, voltage-clamp measurements were performed by using the custom acquisition program GePulse (available at http://users.ge.ibf.cnr.it/pusch/programs-mik.htm) or the program CellWorks (Npi Electronic, Tamm, Germany) and a Turbo TEC-03X or Tec-05X amplifier (Npi Electronic). To estimate the expression level of the constructs varying the osmolarity, different solutions were prepared containing (in mM) 48 NaCl, 2 KCl, 1.8 CaCl₂, 1 MgCl₂, and 10 Heps at pH 7.3 (osmolarity, 120 mOsm), called “Hypo”; “Hypo” was supplemented with mannitol to obtain the “Iso” solution (osmolarity, 200 mOsm) and the “Hyper” solution (osmolarity, 310 mOsm). The investigation of the osmolarity dependence of 8A-VFP/8E-mCherry required additional solutions containing (in mM) 24 NaCl, 2 KCl, 1.8 CaCl₂, 1 MgCl₂, and 10 Heps at pH 7.3 (osmolarity, 74 mOsm), and 7 NaCl, 2 KCl, 1.8 CaCl₂, 1 MgCl₂, and 10 Heps at pH 7.3 (osmolarity, 40 mOsm).

The standard bath solution used for the selectivity measurements contained (in mM) 100 NaCl, 5 MgSO₄, and 10 Heps at pH 7.3 (osmolarity, 215 mOsm). The selectivity to the different ions was estimated substituting sodium chloride with sodium iodide, nitrate, bromide, glutamate, gluconate, bicarbonate, thiocyanate, fluoride, and lactate. Similarly, sodium selectivity was tested by substituting sodium chloride with tetaethylammonium chloride (TEA-Cl). For taurine and glycine, which are neutral at pH 7, we prepared solutions at alkaline pH containing (in mM) 140 taurine, 64 NMDG (N-methyl-D-glucamine), and 5 MgSO₄ at pH 9, and 159 glycine, 38 NMDG, and 5 MgSO₄ at pH 9.3, respectively. Based on previous work (4), the concentration of the negatively charged species was calculated as 78 mM for the taurine solution and 53 mM for the glycine solution assuming pKa values of 8.9 for taurine and 9.6 for glycine, respectively. The selectivity of the LRRC8 constructs was estimated based on the reversal potential change resulting from the replacement of the standard bath solution with the desired solution. Liquid junction potentials were measured and subtracted.

For the selectivity measurements, the reversal potential was determined applying voltages ranging from −60 to 50 mV with 10 mV increments for 50 ms and fitting a parabola to the four points nearest to the zero current potential. The selectivity was estimated according to the Goldman-Hodgkin-Katz equation (21), which for Cl⁻ replacement with different ions became

\[
P_C \times \frac{[\text{Cl}]}{[x]} = e^{\frac{\Delta E_R}{RT}},
\]

where \(P_\text{Cl}\) and \(P_x\) are the permeability of the ion tested and of Cl⁻, [Cl] and [x] are concentrations of Cl⁻ and the x ion, and \(\Delta E_R\) is the shift of the estimated reversal potential. When we replaced NaCl with TEA-Cl, the formula became

\[
P_\text{Na} \times \frac{[\text{Cl}]}{[\text{Na}^+]_{\text{out}}} = \left( e^{\frac{\Delta E_R}{RT}} - 1 \right),
\]

where \(P_\text{Na}\) and \(P_\text{Cl}\) are the permeability of Na⁺ and Cl⁻, \([\text{Cl}]_{\text{out}}\) and \([\text{Na}^+]_{\text{out}}\) are the Cl⁻ concentration inside the oocyte (~30 mM) and the initial external Na⁺ concentration (100 mM), respectively. Here, we assumed that TEA is completely impermeable (compared to sodium).

To estimate LRRCA-mediated currents at different potentials, the “IV-pulse protocol” was applied: a prepulse to −100 mV for 200 ms was followed by voltages ranging from −100 to 60 mV with 20 mV increments for 3000 ms. Pulses ended with a tail to −70 mV for 500 ms.

To study the effect of CBX, we applied increasing concentrations of CBX (10, 20, 50, 100, and 500 μM) dissolved in the “Iso” solution, performing the solution changes when the current was almost at steady state (see Fig. 6A). To quantify the blocking effect of CBX, the steady-state current at various CBX concentrations was estimated by extrapolating to

\[
P_C \times \frac{[\text{Cl}]}{[x]} = e^{\frac{\Delta E_R}{RT}},
\]

where \(P_\text{Cl}\) and \(P_x\) are the permeability of the ion tested and of Cl⁻, [Cl] and [x] are concentrations of Cl⁻ and the x ion, and \(\Delta E_R\) is the shift of the estimated reversal potential. When we replaced NaCl with TEA-Cl, the formula became

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infinite time an exponential function fitted to the time course of the onset of block (see Fig. 6 A). The steady-state current was normalized to the initial current in the “Iso” solution (see Fig. 6 B) and fitted to the Hill equation of the form

\[ p_a = \frac{1 - p_m}{1 + \left( \frac{[CBX]}{K_A} \right)^n + p_m}, \]  

where \( p_a \) is the probability of not being blocked by CBX, \([CBX]\) is the CBX concentration, \( K_A \) the apparent binding constant, \( n \) the Hill coefficient, and \( p_m \) takes into account the leak currents remaining after complete blockage.

**Electrophysiology (patch clamp)**

For patch clamp, the vitelline membrane was mechanically removed by small forceps. To achieve this, oocytes were bathed in a slightly hypertonic medium obtained by mixing a standard shrinking solution (containing (in mM) 200 Na-Aspartate, 20 KCl, 1 MgCl₂, 5 EGTA, and 10 Heps at pH 7.3) with the patch-clamp bath solution (see below) in a 1:2 ratio, resulting in an osmolarity of ~280 mOsm. Oocytes were kept in this solution for at most 2 min.

Single-channel measurements were performed by patch clamp in the cell-attached configuration. The bath solution contained (in mM) 100 N-methyl-D-glucamine-Cl (NMDG-Cl), 2 MgCl₂, 1 EGTA, and 10 Heps at pH 7.3. The extracellular solution (in the pipette) was the same “Iso” solution used for voltage-clamp measurements, containing (in mM) 48 NaCl, 2 KCl, 1.8 CaCl₂, 1 MgCl₂, and 10 Heps at pH 7.3 supplemented with mannitol to obtain an osmolarity of 200 mOsm. Pipettes were pulled from aluminosilicate glass capillaries (Hilgenberg) and had resistances of 3–5 MΩ in the recording solution. To record single-channel events at different potentials, repetitive pulses ranging from 80 to -80 mV with -20 mV increments for 1000 msec were applied in ultrapure water and the pH was adjusted to 7.4 with NaOH. A homemade recording chamber, a hemolysis tube was placed that contained 500 µL of the test solution and 5 µL of D-luciferin and 5 µL of luciferase. Usually, two oocytes were immersed in the tube. Light emitted when ATP reacted with LL was captured by a photomultiplier (R374, Hamamatsu, Hamamatsu City, Japan) fed at high voltage (700–800 V). The resulting signal was amplified in a low-noise amplifier (P16, Grass Valley, Las Cruces, NM), filtered at 10 Hz with a Bessel filter (Frequency Devices, Ottawa, IL), and digitized at 50 Hz using WinWCP (v3.3.3) software (from Professor John Dempster, Strathclyde University, Glasgow, United Kingdom). The amount of ATP released after 7 min was calculated taking into account the signal of standard amounts of ATP in the same experimental conditions. The recording chamber is illustrated in detail in Fig. S1 in the Supporting Material.

**ATP release**

ATP release from hypotonically challenged oocytes was measured using a Luciferase and D-Luciferin mixture (LL). Luciferase extract lanthanum from *Photinus pyralis* (Sigma Aldrich) was resuspended at a concentration of 0.1 µg/µL and desalted in a 10 mL 10 DG column (Biorad, Hercules, CA). D-Luciferin (Sigma Aldrich) was diluted at a concentration of 0.7 µg/µL in ultrapure water and the pH was adjusted to 7.4 with NaOH.

In a homemade recording chamber, a hemolysis tube was placed that contained 500 µL of the test solution and 5 µL of D-luciferin and 5 µL of luciferase. Usually, two oocytes were immersed in the tube. Light emitted when ATP reacted with LL was captured by a photomultiplier (R374, Hamamatsu, Hamamatsu City, Japan) fed at high voltage (700–800 V). The resulting signal was amplified in a low-noise amplifier (P16, Grass Valley, Las Cruces, NM), filtered at 10 Hz with a Bessel filter (Frequency Devices, Ottawa, IL), and digitized at 50 Hz using WinWCP (v3.3.3) software (from Professor John Dempster, Strathclyde University, Glasgow, United Kingdom). The amount of ATP released after 7 min was calculated taking into account the signal of standard amounts of ATP in the same experimental conditions. The recording chamber is illustrated in detail in Fig. S1 in the Supporting Material.

**Uptake measurements**

To measure uptake of [2,2-³H]-taurine, L-[3,4-³H]-glutamate, and [2-³H]-glycine (NEB Radiochemicals, PerkinElmer, Waltham, MA), seven oocytes per individual data point were first incubated for 10 min in Na1X isosmotic solution containing (in mM) 100 NaCl, 2 KCl, 1 CaCl₂, 1 MgCl₂, and 10 Heps/Tris, pH 7.5, or in Na0.5X hypotonic solution (obtained by diluting the Na1X solution 1:1 with water). To initiate uptake, oocytes were transferred into 90 µL Na1X or 90 µL Na0.5X solution supplemented with 100 µM of substrate at 1 µCl/mL, and incubated at 25°C for 20 min. After incubation, the uptake solution was removed, and the oocytes were washed three times in 4 mL ice-cold Na1X solution supplemented with 1 mM substrate. Immediately, each single oocyte was inserted into a scintillation vial and dissolved in 200 µL of 10% sodium dodecyl sulfate, and the radioactivity was counted after the addition of 3 ml of scintillation fluid.

**Water permeability measurements**

Water permeability was assessed by placing individual oocytes into small plates containing hypotonic ND48 medium (ND96Z). This was a two-step procedure in which we first equilibrated the internal osmolality to isosmotic ND48 with mannitol (190 mM), and later placed the oocyte into ND48 (95 mM). Oocyte swelling was recorded by a Nikon CDSS230 stereo-microscope coupled to a Nikon DS-U2 camera. NIS Elements software was used to configure the imaging protocol for data acquisition. Oocyte volume was obtained from the oocyte section area calculated in ImageJ. Water permeability (Pf) was calculated from the initial rate of volume increase with the formula

\[ V_o = \frac{V_o [d(V_o/dt)]}{S_o \times V_o (Osm_{in} - Osm_{out})}, \]

where \( V_o \) is the initial oocyte volume (in cm³), \( S_o \) the initial oocyte area (in cm²), \( V_o \) the molar volume of water (18 cm³/mol), \( Osm_{in} \) the internal oocyte osmolarity (190 mOsm), and \( Osm_{out} \) the ND48 osmolarity.

**Western blot**

Thirty oocytes were homogenized by 20 strokes in an Eppendorf Teflon-glass homogenizer in 10 µL oocyte of buffer containing 1% TX-100, 150 mM NaCl, and phosphate-buffered saline plus protease inhibitors. The homogenate was centrifuged twice at 1000 × g for 10 min at 4°C to eliminate the yolk. The proteins in the supernatant were quantified by the BCA method. Sodium dodecyl sulfate polyacrylamide gel electrophoresis and Western blot experiments were performed as described (22), loading 100 µg of extract. We used the antibodies anti-GFP (ab290, AbCAM, Cambridge, United Kingdom), anti-Cherry (ab125096, AbCAM), and anti-tubulin (MMS-435P, Covance Antibody Products, Princeton, NJ) at 1:100 dilution.
oil-immersion objective. Excitation light at 488 nm from an Argon-Krypton laser (Coherent, Santa Clara, CA) was used to excite the Venus fluorescent protein (VFP) in TIRF mode. Excitation light at 560 nm from a fiber laser (MPB Communications, Montreal, Canada) was used to excite mCherry in TIRF mode. Emission of VFP was collected with a BP 525/50 emission filter (Chroma Technology, Bellows Falls, VT) and that of mCherry was collected with BP 605/52 (Chroma Technology). The fluorescence emission was recorded onto an EMCCD camera (pixel size, 157 nm) (Andor Technology, Belfast, United Kingdom) with an exposure time of 100 ms/frame.

The single-step photobleaching images were achieved by exposing first mCherry to 560 nm laser light at 7.9 W/cm² until all were photobleached, and then, in the same field of view, VFP to 488 nm laser light at 3.2 W/cm² until all were photobleached. For the analysis of the step-photobleaching data, a square region of $3 \times 3$ pixels around the center of each spot was chosen to extract intensity-time traces using custom-written Python software. Background was subtracted locally by considering the average intensity of a $2 \times 2$ pixel region surrounding the molecule of interest. We excluded from our analysis spots that moved by $\gtrsim 1$ pixel before they photobleached, spots that were fluorescent for only one frame, and multiple partially overlapping spots. The signal/noise ratio was $\sim 3$, and it has been shown previously that the counting in step bleaching methods is reliable as long as the signal/noise ratio is 1.5 or above ($23$). The steps in intensity-time traces were counted manually, as previous work ($24$) has indicated that automated fitting ($25$) is not as reliable for this type of data. Photobleaching steps were defined as events in which the mean intensity of the smallest step was greater than at least two standard deviations from the background fluorescence. The intensity of an individual step was determined from traces containing only a single step and was $6.2 \pm 1.6$ a.u. for VFP. This intensity was used as a cut-off to define single steps. Steps that had an intensity greater than two standard deviations away from this mean intensity were likely due to missed events in which multiple fluorescent proteins photobleached simultaneously and these traces were not considered in the analysis. The resulting distribution of the number of steps was compared to either a binomial distribution (Eq. 4) or a Poisson distribution (Eq. 5):

$$q_n = \left( \frac{M!}{n!(M-n)!} \right) P^n (1-P)^{M-n}$$

$$q_n = \frac{\lambda^n e^{-\lambda}}{n!}$$

In Eqs. 4 and 5, $q_n$ is the probability of observing $n$ steps in a time trace. In Eq. 4, $M$ is the total number of subunits and $P$ is the probability that the fluorescent protein will be fluorescent. In Eq. 5, $\lambda$ is the average number of subunits ($26$). The distribution should follow binomial statistics if the stoichiometry of the subunits is fixed and a Poisson distribution if the stoichiometry is free ($23,26$). A possible caveat of the photobleaching method is that incomplete folding efficiency of the fluorescent protein can cause a systematic underestimation of the number of subunits ($27$). We previously showed that VFP folds and matures with high efficiency ($\sim 80\%$) in Xenopus oocytes under similar cytosolic tagging and imaging conditions using the human glycine receptor ($23,24$), and we could reliably determine the pentameric stoichiometry of homomeric $\alpha$-subunit expressed alone, as well as the 3:2 stoichiometry of heteromeric $\alpha/\delta$ complexes ($23$). VFP folding efficiency was independent of the type and oligomeric state of the tagged subunit ($23,24$). Given these results, we relied on the photobleaching of VFP to assess the LRRRC8 subunit stoichiometry.

To determine the density of LRRRC8 subunits per unit area, the total number of fluorescent spots in $40 \mu m \times 40 \mu m$ fields of view (FOVs) was counted manually for mCherry as well as for VFP. A spot was considered if its intensity was within the intensity range of a single photobleaching step. In addition, the number of mCherry and VFP spots that colocalized was counted from those spots that shared the same $3 \times 3$ pixel region in the two channels. Once again, we excluded spots that were fluorescent for only one frame and multiple partially overlapping ones in the same region of interest.

RESULTS

First studies of expression of LRRRC8 proteins in Xenopus oocytes

The recent identification of the members of the LRRRC8 family as the molecular correlates of VRAC ($16,17$) allows the molecular characterization of the proteins underlying these channels. In this work, we took advantage of the fact that enzymatically defolliculated Xenopus oocytes have no endogenous VRAC-like current ($8,28,29$) to perform a characterization of LRRRC8 proteins expressed in these cells. For simplification, we named LRRRC8A as 8A and the other isoforms analogously.

We first verified that collagenase-defolliculated uninjected oocytes do not express endogenous VRAC: both in isotonic conditions (200 mOsm) and under continued perfusion with hypotonic solution (120 mOsm) for 10 min, no significant currents were activated (Fig. 1 A). Injection of 8A or 8E RNA alone did not induce VRAC currents (Fig. 1, B and C). Also, currents in 8A and 8E coinjected oocytes were practically indistinguishable from endogenous currents in isotonic conditions (Fig. 1 D, left). However, in contrast to oocytes expressing 8A or 8E alone (Fig. 1, B and C), currents in 8A and 8E coinjected oocytes slowly activated in hypotonic conditions (Fig. 1 D; Fig. S2 A).

Hypotonicity-induced 8A/8E currents showed outward rectification of the initial current upon the voltage step (instantaneous rectification) and slow inactivation at positive potentials, in agreement with properties found in several mammalian cell lines ($5$). After the activation by hypotonicity, 8A/8E-mediated currents returned slowly to the initial current level upon application of the isotonic solution (Fig. S2 A).

We next coexpressed 8A with the other related homologs, 8B, 8C, and 8D. 8A/8B (Fig. S3 A) and 8A/8C (Fig. S3 B) did not show significant currents either in isotonic conditions or under prolonged exposure to hypotonic solution. Similarly, 8A/8D-induced currents were very small and comparable to those of uninjected oocytes in isotonic conditions. Nevertheless, upon hypotonic stimulation, significant currents could be evoked (Fig. S3 C). However the functional expression level of 8A/8D (Fig. S3 C) was lower compared to 8A/8E (Fig. 1 D).

For the purpose of studying stoichiometry and localization of these heteromers, LRRRC8 proteins C-terminally tagged with VFP or with mCherry (mCh) in the figures) were prepared. Surprisingly, the addition of fluorescent proteins to the C-terminus dramatically increased 8A/8E activity. Indeed, the heteromer 8A-VFP/8E-mCherry exhibited consistent currents even in isotonic conditions (Fig. 1 E)
and with similar kinetics and rectification of untagged volume stimulated 8A/8E-induced currents (Fig. 1 D). Importantly, 8A-VFP/8E-mCherry currents were strongly stimulated by hypotonicity (Fig. 1 E). In these conditions, the current magnitude caused series resistance problems resulting in an only apparent loss of the inactivating kinetics seen in isotonic conditions (Fig. 1 E).

We next tested coexpression of untagged constructs with tagged constructs. 8A/8E-mCherry (Fig. 1 F) and 8A-VFP/8E (Fig. 1 G) resulted in constitutively active currents in isosmotic conditions, which were strongly stimulated by hypotonicity but much smaller than 8A-VFP/8E-mCherry-induced currents (Fig. 1 E). Finally, we tested the expression of 8A-VFP and 8E-mCherry alone. 8A-VFP showed no significant currents even upon hypotonic stimulation (Fig. 1 H). However, 8E-mCherry exhibited small currents that increased upon hypoosmotic stimulation (Fig. 1 I) and after more days of expression (Fig. S4), but remained much smaller (<10%) than those of 8A-VFP/8E-mCherry (compare Fig. 1 I to Fig. 1 E, and see Fig. S4).

Coexpressing 8A fused with mCherry and 8E fused with VFP resulted in activation of currents similar to that for the inverse combination, and also, the coexpression of 8A-VFP with 8E-VFP or of 8A-mCherry with 8E-mCherry activated 8A/8E currents (Fig. S5). On the other hand, adding three copies of the flag epitope did not activate the currents, and adding three copies of the HA epitope led only to a very small activation (Fig. S5). Thus, the size of the added tags appears to be critical for a significant activation, although other factors, such as the charge or the shape of the tag, may also contribute.

We sought to test whether the constitutive activation of currents by adding fluorescent tags was specific for the proteins expressed in oocytes by patch clamping of 8A-VFP/8E-mCherry-transfected human embryonic kidney cells. However, we found this to be extremely difficult because of a very low success rate of giga-seal formation. In the few successful recordings, constitutive VRAC-like channel activity could be observed (data not shown). Because of this difficulty, we concentrated on the oocyte expression system.

**Functional characteristics of the current induced by LRRC8 proteins**

Based on the above described results, we exploited the fluorescently tagged subunits as a useful tool to investigate

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**Figure 1** Functional expression of LRRC8-mediated VRAC currents in *Xenopus* oocytes. (A) Voltage-clamp traces of an un.injected oocyte evoked by the IV-pulse protocol in “Iso” (A, top left) and after 5 min perfusion of “Hypo” solution (A, top right). Mean values of currents at 60 mV from un injected oocytes in “Iso,” and after 5 min and 10 min in “Hypo” solution (A, bottom) (n = 7). (B–G) Currents of single oocytes injected with 8A (B), 8E (C), and coinjected with 8A/8E (D), 8A-VFP/8E-mCherry (E), 8A/8E-mCherry (F), and 8A-VFP/8E (G) in response to the IV-pulse protocol in “Iso” solution and after 5 min perfusion with “Hypo” solution. (B–G, bottom) Current at 60 mV of 8A (B), 8E (C), 8A/8E (D), 8A-VFP/8E-mCherry (E), 8A/8E-mCherry (F), and 8A-VFP/8E (G) in “Iso” and after 5 min and 10 min in “Hypo” solution (n ≥ 4). The dashed line in (A)–(D) indicates an arbitrary threshold of current expression corresponding to 0.5 μA. (H and I) Mean values of currents at 60 mV from oocytes injected with 8A-VFP (H) and 8E-mCherry (I) in “Iso” and after 5 min and 10 min in “Hypo” solution (n ≥ 4). Error bars indicate the standard deviation.
several functional properties of the activity induced by LRRC8 proteins. We prepared mCherry-labeled constructs for the other subunits (B, C, and D) and coexpressed these with 8A-VFP. All constructs were expressed in oocytes as shown by Western blot (Fig. S6), even though 8D-mCherry protein expression was much smaller than that of the other subunits (Fig. S6). The expression level of 8A was similar when expressed alone or coexpressed with the other subunits (Fig. S6A). Differences in molecular weight between the subunits may be due to the variable presence of posttranslational modifications. Interestingly, 8A-VFP/8B-mCherry showed no significant currents even after 10 min of hypotonic stimulation (Fig. 2A), although both subunits were expressed (Fig. S6). 8A-VFP/8C-mCherry resulted in large constitutive currents in isotonic conditions (Fig. 2B), comparable to those for 8A-VFP/8E-mCherry (Fig. 2D). Interestingly, 8A-VFP/8C-mCherry-induced currents exhibit no time-dependent inactivation at positive potentials (Fig. 2B). 8A-VFP/8D-mCherry co-injection resulted in smaller constitutive and volume-stimulated currents compared to 8A-VFP/8E-mCherry (Fig. 2, C and D). At positive voltages, 8A-VFP/8D-mCherry (Fig. 2C) displayed time-dependent inactivation kinetics similar to those of 8A-VFP/8E-mCherry (Fig. 2D).

Taking advantage of the stable constitutive current exhibited by fluorescently tagged LRRC8 proteins in isotonic conditions, we could establish the selectivity sequence for various subunit combinations using reversal potential measurements as illustrated in Fig. S7. The permeability sequence found for 8A-VFP/8E-mCherry was SCN$^-$ > I$^-$ > NO$_3^-$ > Br$^-$ > F$^-$ > HCO$_3^-$ > glycine > taurine > lactate > aspartate > glutamate > glutamate > Na$^+$. This is very similar to that previously found for VRACs (3) with a larger permeability to iodide than to chloride (Fig. 2E, left). As already reported, this is an Eisenman anion sequence I (I$^-$ > Br$^-$ > Cl$^-$ > F$^-$) corresponding to an anion binding site of weak field strength (3,6). 8A-VFP/8E-mCherry was instead substantially impermeable to Na$^+$ ($P_{Na}/P_{Cl}$ ~2%), confirming that LRRC8 proteins are anion channels (Fig. 2E, left). A very similar, but not identical, permeability sequence was found for 8A-VFP/8C-mCherry: SCN$^-$ > I$^-$ > Br$^-$ > NO$_3^-$ > F$^-$ > glycine > HCO$_3^-$ > taurine > lactate > aspartate > glutamate = glutamate > Na$^+$ (Fig. 2E, middle). Interestingly, the permeability sequence of 8A-VFP/8D-mCherry...
was SCN− > I− > glycine = NO3− > Br− > taurine > F− > HCO3− = aspartate > lactate = glutamate > glucoc- 
nate > Na+ (Fig. 2 E, right; Fig. S7 C). Thus, unlike 8C 
and 8E, the 8D-mCherry subunit confers a very large perme- 
ability to glycin (Pgly/PCl ~1.17) and taurine (Ptau/PCl 
~0.76), suggesting that 8A-VFP/8D-mCherry is more effi-
cient at transporting these substrates than the other hetero-
mers, in agreement with recent published data (19).

Based on the large functional expression of 8A-VFP/8E-
mCherry, we chose this heteromer to study single-channel 
properties of LRRC8-mediated currents. Single-channel 
measurements were performed on 8A-VFP/8E-mCherry-in-
jected oocytes in the cell-attached configuration monitoring 
channel activity at potentials ranging from 80 to −80 mV. 
Typical recordings are shown in Fig. 3 A. In agreement 
with previous work on endogenous VRAC (5–7) and recent 
studies addressing LRRC8 reconstituted in bilayers (20), 
we found a pronounced outward rectification of the single-
channel current-voltage relationship (Fig. 3, A and B), 
explaining the outward rectification of the macroscopic cur-
rents. At 80 mV, we measured a single-channel current i = 
2.57 ± 0.29 pA (chord conductance = 32 pS) (Fig. 3 C); at 
20 mV, we found i = 0.21 ± 0.04 pA (chord conductance = 
11 pS); the conductance further decreased at negative poten-
tials with i = −0.37 ± 0.06 pA (chord conductance = 6 pS) 
at −60 mV (Fig. 3 B), which is very similar to what has 
been recently described (20), strongly indicating that the 
observed single-channel currents are mediated by LRRC8 
proteins. Interestingly, the channels showed flickery open-
ings at all voltages (Fig. 3 A).

**Flux of organic osmolytes and ATP through 
LRRC8 proteins**

Reversal potential measurements (Fig. 2 E) indicated that 
the expression of LRRC8 proteins induces a conductance 
that is permeable to amino acids such as glutamate and 
glycine as well as amino acid derivatives such as taurine.

To obtain additional independent evidence that these 
compounds can pass through LRRC8 proteins, we per-
formed taurine, glutamate, and glycine radiotracer experi-
ments. As the endogenous permeability for taurine and 
glutamate is very low (30,31), loading these substrates for 
efflux experiments would be rather difficult. However, we 
reasoned that influx experiments, although probably not in 
linear conditions, were feasible. We thus expressed 8A/8E 
and 8A-VFP/8E-mCherry and measured the uptake of radio-
labeled taurine and glutamate after incubation in an isotonic 
or a hypotonic solution. In isotonic conditions, compared 
with uninjected oocytes, higher influx of radioactively 
labeled taurine (Fig. 4 A) and glutamate (Fig. 4 B) was 
observed in oocytes coexpressing untagged 8A and 8E, 
and it further increased in oocytes coexpressing 8A-VFP 
and 8E-mCherry. After incubation in a hypotonic solu-
tion, both groups showed high amounts of labeled taurine 
(Fig. 4 A) and glutamate (Fig. 4 B). We analyzed the 
influx of taurine, glutamate, and glycine in isotonic (I) or 
hypotonic conditions (H) of 8A-VFP coexpressed with 
all the different subunits (8B, 8C, 8D, and 8E) tagged 
with mCherry. All heteromers except 8A-VFP/8B-mCherry 
showed influx of taurine (Fig. 4 C) and glutamate (Fig. 4 D). 
Also, glycine influx was significantly increased in tagged

**FIGURE 3** Single-channel studies of the 8A-VFP/8E-mCherry heteromer. 
(A) Representative traces from a cell-attached patch at 80, 40, and −80 mV. 
(B) Single-channel current-voltage relationship. Mean single-channel cur-
rents are plotted versus the corresponding potentials ranging between 
80 and −80 mV. Note the outward rectification (n ≥ 4) For −80, −40, and 
−20 mV, n = 2). Error bars indicate the standard deviation. (C) Amplitude 
histogram of the recording at 80 mV in control conditions. The dashed line 
represents the fit with a sum of three Gaussians. The dominant current level 
has an amplitude of 2.5 pA. To see this figure in color, go online.
8A/8C-, 8A/8D-, and 8A/8E-coexpressing, but not 8A/8B-coexpressing oocytes in isotonic and even more in hypotonic conditions (Fig. 4E). However, the difference to noninjected oocytes was smaller compared to that seen for taurine and glutamate influx (Fig. 4, C–E).

The release of osmolytes, especially taurine, through the VRAC channel is important for the process of RVD induced by swelling (4). Xenopus oocytes show low water permeability, reflecting the lack of aquaporins and the lipid composition of the membrane (32). Water permeability can be dramatically increased by the expression of aquaporin-1 (AQP1) such that hypotonic exposure leads to oocyte swelling and membrane disruption (32). To test whether LRRC8 proteins are able to protect oocytes from such cell swelling, we coexpressed AQP1 with 8A-VFP/8E-mCherry. Indeed, swelling was dramatically reduced and was even lower than that seen in oocytes without AQP1 (Fig. 4F). To rule out that RNA competition nonspecifically reduced AQP1 expression in coinjected oocytes, we coinjected AQP1 with 8A-VFP co-expressed with 8B-mCherry, 8C-mCherry, 8D-mCherry, and 8E-mCherry (n ≥ 3 for all co-expressions) in Iso “I” and Hypo “H” solution. (F) Time course of normalized oocyte volume in ND48 solution and apparent water permeability mean values (inset) of uninjected oocytes (n = 6), AQP1 + 8A-VFP (n = 8), and AQP1 + 8A-VFP/8E-mCherry (n = 10) injected oocytes. Data indicate the mean ± SE. *p < 0.05, **p < 0.01, ***p < 0.001.
ATP release strongly depended on osmolarity (Fig. 5 C) and was blocked by CBX (Fig. 5 D). Finally, we analyzed ATP release in oocytes coexpressing 8A-VFP with other mCherry-tagged subunits (B, C, and D). A much smaller but significant ATP release was observed for 8A-VFP/8C-mCherry, whereas it was not statistically significant in heteromers containing tagged 8B or 8D (Fig. 5 D, inset). These results suggest that ATP permeates better through 8A/8E channels than through 8A/8C channels. We thus tested whether ATP differently affects the currents mediated by these subunit combinations. However, addition of 1 mM ATP blocked by ~50% both 8A-VFP/8E-mCherry- and 8A-VFP/8C-mCherry-mediated currents (data not shown).

Mechanism of CBX inhibition of LRRC8-mediated currents

We next aimed to characterize the mechanism of CBX inhibition of LRRC8-mediated currents. CBX is the succinyl ester of glycyrrhizic acid that was shown previously to inhibit VRAC currents (10) and recently also LRRC8-mediated currents (17). Therefore, we tested CBX on 8A-VFP/8E-mCherry-induced currents at various concentrations from 10 to 500 μM (Fig. 6 A). Increasing [CBX] resulted in increasing block of the currents. For low [CBX], complete steady-state inhibition was difficult to achieve in reasonable recording times (Fig. 6 A). Thus, to estimate steady-state block, we fitted an exponential function to the time course of the onset of block and extrapolated it to infinite time (Fig. 6 A, solid lines). Using this procedure, we found ~59% residual current at 10 μM CBX and only ~7% at 100 μM CBX. The residual currents at 500 μM CBX were at the level of endogenous currents, indicating that block is practically complete at high [CBX]. We fitted the CBX block by a simple binding curve (see Materials and Methods). The best fit yields an apparent affinity constant (K_A) of 11.9 μM and a Hill coefficient (n) of 1.8 (Fig. 6 B, red line). The Hill coefficient of 1.8 suggests that more than one CBX molecule is required to block the channel. In fact, fixing the Hill coefficient at 1 yields a poor fit to the data (Fig. 6 B, black dashed line). However, since the experiments were performed without washout control, further evidence is needed to draw a firm conclusion on the number of CBX binding sites.
Bioinformatic analyses have revealed a phylogenetic relation of LRRC8 proteins to pannexins (34). Experimental evidence based on cross-linking and native gels suggests that pannexins may form hexameric complexes (35), although other studies suggest that pannexin-2 may assemble into heptamers or octamers (36). Recent cross-linking studies suggest that also, LRRC8 complexes may contain six to eight subunits (20). However, pannexins are homomeric (37), whereas LRRC8 channels are multimERIC complexes of LRRC8A and other LRRC8 members (17).

To obtain insight into the stoichiometry of LRRC8 channels, we used TIRF microscopy to visualize the fluorescently tagged subunits in the membrane at single-particle resolution. When 8A-VFP was injected alone, we observed a variable but overall large number of diffraction-limited spots (150 ± 30 spots/field of view (FOV), n = 3 FOVs). For 8E-mCherry alone, the number of spots was much smaller (21 ± 13 spots/FOV, n = 4). Upon coexpression of 8A-VFP with 8E-mCherry, the number of 8E-mCherry spots significantly increased (42 ± 16 spots/FOV, n = 11, p = 0.004), whereas the number of 8A-VFP spots slightly decreased (83 ± 24 spots/FOV, n = 11). Twenty-six percent of the 8A-VFP spots colocalized with 8E-mCherry spots, and 51% of 8E-mCherry spots colocalized with the 8A-VFP spots (n = 11, p = 0.003) (Fig. S7 A). These results suggest that 8A and 8E form oligomERIC complexes at the plasma membrane, that 8E relies on 8A to achieve efficient plasma membrane expression, as previously reported (17), and that 8A levels at the plasma membrane are independent of 8E.

To investigate whether more than two different subunits may be associated in the same complex, we coexpressed untagged 8A with 8E-VFP and 8D-mCherry and detected colocalization of 8E-VFP and 8D-mCherry (Fig. S8 A). Assuming that 8A is obligatorily present, this indicates that LRRC8 complexes may be formed by more than two different subunits.

Since the tagged subunits were relatively immobile in the oocyte membrane, we used single-step photobleaching to count and estimate the number of subunits within the hetero-oligomers. This method has been used to determine the stoichiometry of small oligomers (five to six subunits) (24,38). For an oligomer with fixed stoichiometry, the distribution of the observed number of photobleaching steps is binomial, whereas for a variable stoichiometry the distribution is expected to be Poissonian (see Materials and Methods).

We extracted intensity-time traces from 8A-VFP particles that colocalized with 8E-mCherry particles and thus were heteromERIC (Fig. 7 B). The number of photobleaching steps showed a broad distribution (Fig. 7 C) and we could reliably count only up to six photobleaching steps, since for more than six steps, the probability of multiple fluorescent proteins photobleaching simultaneously increases (Fig. S8 B). This broad distribution is not compatible with a binomial distribution assuming 80% probability for VFP to be fluorescent and a fixed stoichiometry ranging from 1 to 6.
Instead, it fits well to a Poisson distribution (Fig. 7 C), indicating that the number of 8A subunits present in LRRC8 hetero-oligomers is variable. We calculated an average number of ~3 8A-subunits within the hetero-oligomers. As this number is very similar to what has been observed recently in cross-linking experiments (20), we assume that we are predominantly measuring fluorescence from a single VRAC channel and not from clusters of channels.

To similarly determine the number of 8E subunits in the hetero-oligomers, we injected an equimolar ratio of 8E-VFP and 8A-mCherry and extracted intensity-time traces from 8E-VFP spots that colocalized with 8A-mCherry spots (Fig. 7 B). The expression level of the two subunits was similar to that of the 8A-VFP/8E-mCherry combination (70 ± 22 8A-mCherry spots and 49 ± 27 8E-VFP spots/FOV, n = 9). In addition, the percentage of colocalization between 8A-mCherry and 8E-VFP was similar (28% of 8A-mCherry colocalized with 8E-VFP and 38% of 8E-VFP colocalized with 8A-mCherry; n = 9). As for 8A-VFP (Fig. 7 C), we observed a broad distribution for the number of photobleaching steps that fits well with a Poisson distribution (Eq. 5). Taken together, these results suggest that the number of 8A and 8E subunits in LRCC8 hetero-oligomers is variable and that on average the total number of subunits is >5. We note that since the folding of VFP is incomplete but high (~80%), the number of subunits may be slightly underestimated.

It has been reported that overexpression of LRRC8A decreased VRAC currents in a manner similar to that...
observed in the inhibition by RNA interference (16,17). Given that in *Xenopus* oocytes it is very easy to change the relative expression of each subunit by changing the amount of injected cRNA, we compared the currents induced by injection of equimolar amounts of 8A-VFP and 8E-mCherry (1:1) with those induced by increasing the relative amount of 8A-VFP versus 8E-mCherry (3:1) (Fig. 8C). Increasing the relative amount of 8A-VFP decreased the currents by ~75–80% (Fig. 8C, left), with similar results for 8A-mCherry and 8E-VFP (Fig. 8C, right). We also counted the number of photobleaching steps for 8A-VFP when the ratio was 3:1 and found an average number of ~4 A-subunits in the complex, indicating that the stoichiometry of each LRRC8 protein is not fixed but likely depends on their relative abundance (Fig. 8D).

**Fluorescent-tagged LRRC8 proteins show a shift in their osmolarity dependence**

Physiologically, VRAC is closed under isotonic conditions, and indeed, oocytes injected with wild-type 8A/8E or 8A/8D subunits do not show significant currents in isotonic conditions (Fig. 1; Fig. S1). Thus, the constitutive currents induced by the fluorescently tagged LRRC8 proteins in isosmotic conditions raise the question of whether the tags render the channels constitutively open or they are still able to close. To address this question, we studied in quantitative detail the dependence of 8A-VFP/8E-mCherry-induced currents on the extracellular osmolarity.

Interestingly, the constitutively active currents seen after expression of the fluorescently tagged subunits are reduced by application of a hyperosmotic solution (310 mOsm) to <20% (Fig. 8). This result indicates that the addition of the fluorescent tags leads to a “shift” of the osmosensitivity of the channels such that a basal activation is present even under isotonic conditions. Decreasing the osmolarity to 120 mOsm leads to a ninefold activation of 8A-VFP/8E-mCherry-mediated currents compared to the isotonic solution, whereas further osmolarity reductions (to 74 and 40 mOsm) result in 17-fold and 25-fold current increases, respectively (Fig. 8). Thus, the osmolarity dependence of these channels is not even saturated at 40 mOsm (Fig. 8, right). Because the untagged constructs responded more slowly to hypotonic challenges, it was unfeasible to assay their osmosensitivity in a similar manner.

**DISCUSSION**

VRAC currents mediate RVD in many cells and thus play an important physiological role (3). After a long search for the molecular correlates of VRACs (15), finally two independent groups identified LRRC8 proteins as major constituents of the VRAC channel (16,17). However, even after the successful identification, the investigation of the properties of the channel remained challenging, requiring knock-out cell lines for heterologous expression (17,18,20). Here, we show that a simple approach (C-terminal tagging by fluorescent proteins) resulted in constitutive open channels of large magnitude. This allowed investigation of the functional properties of the currents induced by LRRC8 proteins in *Xenopus* oocytes. Using this system, we could confirm previous suggestions that VRAC is able to create a path for many anions and osmolytes, including taurine, glutamate, glycine, and ATP (3,5,39). We could detect significant ATP release for the heteromers 8A-VFP/8E-mCherry and 8A-VFP/8C-mCherry. ATP release was not detected for 8A-VFP/8D-mCherry, probably due to the lower expression of this heteromer in the oocyte system. Expression of LRRC8A alone together with AQP1 has been previously shown to induce chloride currents in oocytes (40). However, we believe that these currents could be due to indirect effects on endogenous currents, as they were also observed when expressing the unrelated membrane protein anoctamin 10 together with AQP1 (40). In our study, LRRC8A alone never gave rise to any significant currents.

Several pieces of evidence support the conclusion that the activity measured in our experiments is due to the expression of the exogenous proteins. 1) As already described in cell lines (17–19), we observed differences in inactivation and selectivity properties for different LRRC8 heteromers. Moreover, single-channel analyses at different voltages were in agreement with previous studies of the VRAC current (3,7) and what has been observed in bilayers after LRRC8 reconstitution (20). 2) We could observe flux of radioactive taurine, glutamate, and glycine, whereas un.injected oocytes have negligible flux for taurine and glutamate and less flux for glycine. 3) Currents were blocked by CBX (10). It has to be considered that VRAC currents have been previously observed in manually isolated *Xenopus* oocytes, but never in collagenase-treated oocytes, as we also confirmed (8,29). It has been suggested that the
VRAC channel might be present in the follicle cells but not in the oocyte membrane (28). We envisage that the endogenous LRRC8 proteins are kept in a dormant state, as has been found in other channels or transporters (41,42). Only by injecting large amounts of LRRC8E-mCherry RNA and after several days could we observe VRAC-like currents. However, compared to the current induced by coexpressing LRRC8A-VFP with LRRC8E-mCherry, the current induced by LRRC8E-mCherry alone was very small (<10%). Thus, a possible endogenous background can be rather safely neglected.

Phylogenetic analyses suggested that the LRRC8 protein family evolved from an ancestral pannexin protein in chordates (34). This homology can clearly be observed via the conservation of cysteine residues in extracellular loops, which are also present in connexins (43). Recently, the first extracellular loop of pannexin1 was found to be involved in CBX binding (13), in agreement with our finding that CBX blocks LRRC8-mediated currents acting extracellularly. This evolutionary relationship also suggested that LRRC8 proteins might form hexameric complexes (37). Our results using single-step photobleaching suggest that the total number of subunits (LRRC8A plus LRRC8E) on average is >5. Since the method relies on counting inactivation steps of VFP, it is challenging to estimate stoichiometry when the number of subunits is very high. However, we found that, when expressed at equimolar levels, the average numbers of LRRC8A and LRRC8E subunits are similar. It will be interesting to find out whether LRRC8A and LRRC8E subunits occupy segregated parts of the heteromer or are interdigitated. Our data further suggest that the complex might contain more than two different LRRC8 subunits. In addition, we found that the stoichiometry is likely not fixed, since the distribution of the number of bleaching steps did not follow a binomial distribution and because the distribution changed upon altering the relative amount of each subunit. Thus, we hypothesize that the dominant negative functional effects observed by the overexpression of LRRC8A may be caused by the formation of nonfunctional stoichiometries. Similar dominant negative effects are observed when overexpressing nicotinic receptors (44), probably by having complexes of nonnative stoichiometries. Thus, we envisage that changing the relative proportion of each subunit may be a way to regulate VRAC channel activity. In this regard, it is interesting to note that heterologous coexpression of LRRC8A and LRRC8B subunits does not result in functional activity (this work and (17)), although the proteins are located at the plasma membrane (17). However, cells expressing only endogenous LRRC8A and LRRC8B (knock-out of LRCC8C/D/E) exhibit VRAC currents (17). It may be that heteromers containing LRRC8B need a stimulus to be active, or, alternatively, LRRC8B could work as an inhibitory subunit.

Many different mechanisms, such as oxidation, phosphorylation, ionic strength, intracellular Na\(^+\) concentration, changes in subcellular localization, and membrane or cytoskeleton stretch (3,6,14,15,20,45–47), have been proposed to regulate the activity of VRAC channels. Here, we found that the addition of fluorescent proteins to the C-terminus, which contains the leucine-rich repeats (48), resulted in constitutively active channels, which are, however, further activated by hypotonicity and inhibited by hypertonicity. We thus suggest that some of the mechanisms known to regulate channel activity may influence the conformation of the C-terminus. In a similar manner, one of the proposed mechanisms for pannexin-1 activation involves the C-terminus, which can be cleaved by caspase, and which could act as a plug in the transmembrane pore (49–52). Further studies are needed to determine whether this mechanism is also occurring in LRRC8 proteins.

In summary, our work established the Xenopus oocyte system as a valid expression system for LRRC8 proteins, and it describes, to our knowledge, new characteristics of this channel. We consider the oocyte system as a complementary approach to the expression in knockout cell lines (17). The oocyte system offers the advantage of the possibility of performing a variety of electrophysiological, optical, and biochemical assays on single cells (53–55) with the ability to control the expression of each subunit independently. Furthermore, the constitutive channel activity in physiological conditions observed after C-terminal fusion of fluorescent proteins in oocytes could facilitate drug screening of this channel. We envisage that the easiness of the oocyte system described here, combined with the activation conferred by the addition of the fluorescent proteins, could be crucial to discover novel structure-function relationships of the VRAC channel formed by LRRC8 proteins. We believe that our system also will be helpful in the unraveling of molecular determinants of novel physiological and pathophysiological roles of LRRC8 proteins, as, for example, their involvement in chemotherapeutic drug resistance (19).

SUPPORTING MATERIAL

Eight figures are available at http://www.biophysj.org/biophysj/ supplemental/S0006-3495(16)30752-4.

AUTHOR CONTRIBUTIONS

The present project was initiated at the lab of R.E. Functional 2-EV electrophysiological measurements were done in the lab of M.P. and R.E. by A.G. and H.G.-P., and single-channel studies were done by A.G. and M.P.H.G.-P., L.L.-C., R.E., and M.L. performed TIRF and step-bleaching experiments and analyses, H.G.-P., R.E., and C.S. performed ATP experiments. H.G.-P. and R.E. performed radioactive uptake and volume measurement experiments. V.F.-D and F.C. helped in the design of experiments. All authors planned and analyzed experiments. R.E. and M.P. supervised experiments. R.E. and M.P were the primary writers and managed production of the manuscript with critical input from all the authors, who read and provided feedback on the manuscript.
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