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Quantifying Ca^{2+} Current and Permeability in ATP-gated P2X7 Receptors*

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Xin Liang (梁欣)[‡], Damien S. K. Samways[§], Kyle Wolf[¶], Elizabeth A. Bowles[‡], Jennifer P. Richards[‡], Jonathan Bruno[¶], Sébastien Dutertre^{**}, Richard J. DiPaolo[¶], and Terrance M. Egan^{‡1}

From the [‡]Department of Pharmacological and Physiological Science and Center for Neuroscience, and the Departments of [¶]Molecular Microbiology and Immunology and ^{¶¶}Internal Medicine, Saint Louis University School of Medicine, St. Louis, Missouri 63104, the [§]Department of Biology, Clarkson University, Potsdam, New York 13699, and the ^{**}Institut des Biomolécules Max Mousseron, UMR 5247, Université Montpellier 2, CNRS, Montpellier, France

Background: Ca^{2+} triggers many of the actions of extracellular ATP.

Results: We measured the Ca^{2+} component of the ATP-gated non-selective cation current of P2X7 receptors.

Conclusion: Ca^{2+} flux varied with the species of origin, the splice variant, and the agonist concentration.

Significance: Our results suggest that domains that lie outside of the pore regulate the Ca^{2+} flux of P2X7 receptors.

ATP-gated P2X7 receptors are prominently expressed in inflammatory cells and play a key role in the immune response. A major consequence of receptor activation is the regulated influx of Ca^{2+} through the self-contained cation non-selective channel. Although the physiological importance of the resulting rise in intracellular Ca^{2+} is universally acknowledged, the biophysics of the Ca^{2+} flux responsible for the effects are poorly understood, largely because traditional methods of measuring Ca^{2+} permeability are difficult to apply to P2X7 receptors. Here we use an alternative approach, called dye-overload patch-clamp photometry, to quantify the agonist-gated Ca^{2+} flux of recombinant P2X7 receptors of dog, guinea pig, human, monkey, mouse, rat, and zebrafish. We find that the magnitude of the Ca^{2+} component of the ATP-gated current depends on the species of origin, the splice variant, and the concentration of the purinergic agonist. We also measured a significant contribution of Ca^{2+} to the agonist-gated current of the native P2X7Rs of mouse and human immune cells. Our results provide cross-species quantitative measures of the Ca^{2+} current of the P2X7 receptor for the first time, and suggest that the cytoplasmic N terminus plays a meaningful role in regulating the flow of Ca^{2+} through the channel.

All P2X receptors transduce a significant Ca^{2+} flux at the resting membrane potential (1) that triggers many of the physiological and pathophysiological actions of extracellular ATP (2–6). The molecular physiology of the Ca^{2+} flux is poorly understood despite outstanding recent advances in functional and structural studies (7). This is particularly true for the P2X7 receptor (P2X7R) that is unusually sensitive to

allosteric block by extracellular Ca^{2+} (8), a fact that makes characterization of the Ca^{2+} component of the ATP-gated current particularly problematic (9, 10).

P2X receptors are a family of seven ATP-gated ion channels (P2X1R–P2X7R) that subserve a diverse range of functions in a subunit selective manner. The prominent expression of P2X7Rs in lymphocytes, macrophages, and microglia (11–14) suggests a vital role for this subtype in the immune response (15). P2X7Rs are inactive in healthy tissue because limited release and rapid hydrolysis keep the concentration of extracellular ATP ($[\text{ATP}]_o$) low (<100 nM) (16–18). However, the $[\text{ATP}]_o$ rises to millimolar concentrations at sites of stress, cellular injury, tumor necrosis, and phagocytic degranulation (16, 18–21) where it acts as a “danger” signal that initiates innate immunity by stimulating the caspase-1-activating platform known as the “inflammasome” (22–24). The result is an increased production and rapid release of pro-inflammatory cytokines by mononuclear phagocytes (15, 24–26), ultimately leading to greater inflammation and/or cell death (13, 15). In many cases, the inward flow of Ca^{2+} through the P2X7R is a key component of this response (28–30). Furthermore, the sustained rise in the intracellular concentration of free Ca^{2+} that follows activation of P2X7Rs directly triggers macrophage apoptosis independent of pro-inflammatory signaling cascades (31).

P2X7Rs are also expressed by neurons and glia of the peripheral and central nervous systems (5) where they influence differentiation, homeostasis, and disease in part by increasing intracellular Ca^{2+} (25, 32–36). In the peripheral nervous system, the elevated $[\text{ATP}]_o$ that accompanies inflammatory bowel disease leads to rapid degeneration of gut neurons by direct activation of the P2X7Rs of the myenteric plexus (37). In the eye, subretinal hemorrhage increases the $[\text{ATP}]_o$, leading to a P2X7R-dependent Ca^{2+} influx that activates caspase-8 and initiates photoreceptor cell apoptosis (38). In the central nervous system, P2X7Rs are found on microglia, oligodendrocytes, and astrocytes, where they contribute to the genesis of neuropathic and chronic inflammatory pain (25) and influence cell survival (39). Although the presence of P2X7Rs on central neu-

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This paper is dedicated to the memory of Thomas H. Steinberg, M.D. (1952–2013), who was a true pioneer in the study of ATP-gated P2X7Rs.

¹ To whom correspondence should be addressed: 1402 South Grand Blvd., St. Louis, MO 63104. Tel.: 314-977-6429; Fax: 314-977-6411; E-mail: egantm@slu.edu.

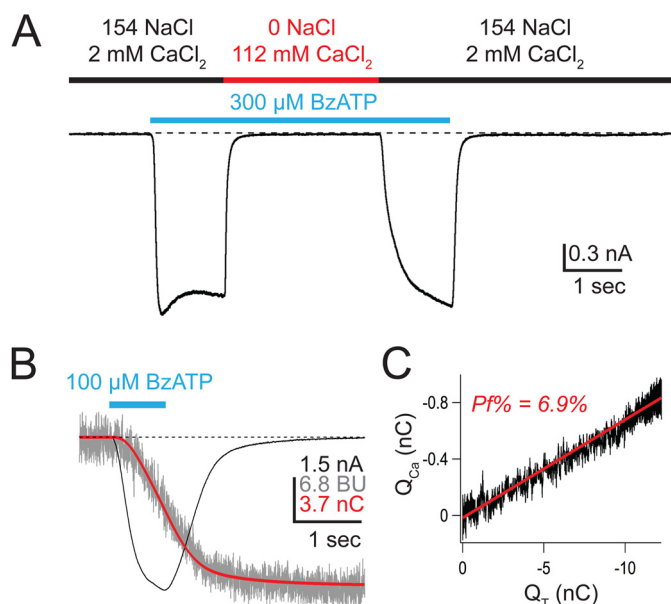


FIGURE 1. **Measuring the Ca²⁺ current of rat P2X7aR.** *A*, the inward current evoked by a supra-maximal concentration ($EC_{50} = 20 \mu\text{M}$ (87)) of BzATP (cyan bar) is blocked in high extracellular Ca²⁺. BzATP was applied first in an extracellular solution containing predominately NaCl (black bar), followed immediately by an extracellular solution containing predominately CaCl₂ (red bar). The high $[Ca^{2+}]_o$ completely blocks the BzATP-gated current. The current quickly recovers upon reintroduction of the NaCl solution. *B*, shows the BzATP-gated membrane current (black trace, in nA), the change in F_{380} (gray trace, in bead units), and the integrated current (red trace, equal to Q_T , in nC). *C*, Q_{Ca} (calculated from F_{380} ; see "Experimental Procedures") is plotted against Q_T . The result is a straight line (slope = Pf), demonstrating that the change in fluorescence results entirely from Ca²⁺ entering through the channel (50). BU, bead units.

rons is controversial (6, 40), recent data suggest they play a role in modulation of release of transmitters and hormones (41, 42). For example, ATP-gated Ca²⁺ influx through presynaptic P2X7Rs triggers glutamate release from cortical synaptosomes (43, 44), GABA release from hippocampal slices (45), and arginine-vasopressin release from neurohypophysial terminals (46).

In this paper, we provide a quantitative measure of the Ca²⁺ component of the P2X7 current. The usual means of achieving this goal is to determine the permeability of Ca²⁺ relative to a reference monovalent cation (47). Unfortunately, measurement of relative Ca²⁺ permeability typically requires a high concentration (10–112 mM) of extracellular Ca²⁺ ($[Ca^{2+}]_o$) that blocks the rat P2X7 current (Fig. 1A) with an IC_{50} of ~3 mM (9, 48). To circumvent this problem, we used the dye-overload method to measure the percent fractional Ca²⁺ current ($Pf\%$)² of the ATP-gated response determined in the presence of less than 2 mM free $[Ca^{2+}]_o$ (49, 50). The $Pf\%$ method has distinct advantages over traditional reversal potential measurements of permeability; it is a direct and model-independent measure of Ca²⁺ entry obtained using physiological concentrations of extracellular ions (51–53). By contrast, reversal-based techniques do not measure the physiologically relevant Ca²⁺ current, require high concentrations of extracellular Ca²⁺, and make the tenuous assumption that the Ca²⁺ current follows the assumptions of the Goldman-Hodgkin-Katz equation (54, 55).

² The abbreviations used are: $Pf\%$, percent fractional Ca²⁺ current; BzATP, benzoyl-ATP; NMDG, *N*-methyl-D-glucamine.

We measured the $Pf\%$ of an array of recombinant and native P2X7Rs, and found an unexpected variability in the contribution of Ca²⁺ to the total ATP-gated membrane current. The variability was most pronounced in the zebrafish P2X7R and in splice variants of murine P2X7Rs. Ten splice variants (P2X7bR–P2X7kR) have been identified from a range of species since the initial molecular cloning of the first P2X7R from rat (now called rat P2X7aR) (6). Only two of these (human P2X7bR and murine P2X7kRs) form functional homomeric receptors (56, 57). We discovered that mouse and rat splice variants with identical pore-lining sequences show significantly different $Pf\%$, suggesting that domains that lie outside of the channel pore regulate the Ca²⁺ component of the ATP-gated current.

EXPERIMENTAL PROCEDURES

Human and animal procedures were approved by the Institutional Review Board and the Institutional Animal Care and Use Committee, respectively, of the St. Louis University School of Medicine.

Isolation of Human Macrophages—Monocyte-derived macrophages were obtained as described by Norenberg *et al.* (58). Blood obtained from healthy volunteers was centrifuged at $500 \times g$ for 10 min to isolate plasma, buffy coat layer, and erythrocytes. The buffy coat layer was removed, diluted 2:1 with cold physiological buffered saline (PBS) with no added divalent cations, and then overlaid on 15 ml of Histopaque 1077 (Sigma) in a 50-ml centrifuge tube. The tube was centrifuged at $900 \times g$ for 30 min to produce an interfacial layer of mononuclear cells, and platelets were isolated and then processed through three PBS wash and spin ($250 \times g$ for 7 min) cycles. After the final spin, the pelleted cells were resuspended in 8 ml of a culture medium made of SensiCell™ RPMI 1640, 7.5% heat-inactivated autologous plasma, 100 units/liter of penicillin, and 100 μg/ml of streptomycin (all from Life Technologies). The cell suspension was then divided equally among eight 4-well plates previously loaded with circular 13-mm glass coverslips (Gold Seal Cover Glass, Thermo Scientific, Waltham, MA), and placed in a humidified 5% CO₂ incubator for 2 h. Subsequently, the plates were washed several times with warm PBS to remove non-adherent cells, and the remaining cells were cultured in 0.5 ml of the culture medium for 6–14 days. Lipopolysaccharide (1 μg/ml, Sigma) was added for 6–12 h immediately preceding the start of the experiments.

Isolation of Mouse Macrophages—Peritoneal macrophages were obtained with modifications as described by Davies and Gordon (59). Two or three adult mice were intraperitoneally injected with 1 ml of sterile 3% Brewer's thioglycolate (Thermo Scientific) using a 25-gauge needle. Four days later they were sacrificed via carbon dioxide inhalation and cervical dislocation. The abdominal skin was soaked with 70% ethanol, pulled up with sterile forceps, and cut to expose the peritoneal cavity. The peritoneum was injected with an ice-cold high-glucose Dulbecco's modified Eagle's medium (DMEM; with added glutamine and sodium pyruvate) (Life Technologies) using a 20-gauge needle, and the mouse was vigorously shaken for a few seconds. The suspended cells were then harvested by removing the DMEM from the peritoneal cavity using a 25-gauge needle. The cell suspensions obtained from individual mice were

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pooled, and the cells were pelleted by centrifugation (5 min at 200 × g). The cell pellet was resuspended in 5 ml of cold Red Cell Lysis Buffer (Sigma), incubated for 5 min at room temperature, and then repelleted. The supernatant was discarded, and the remaining cells were resuspended in DMEM, 10% heat inactivated fetal bovine serum (FBS; Thermo Scientific), and non-essential amino acids (Sigma). The cells were plated at a density of 500,000 cells/35-mm tissue culture dish and cultured in a humidified 5% CO₂ incubator for up to 14 days. Two days before the experiment, the cells were replated onto 13-mm glass coverslips. Lipopolysaccharide (1 μg/ml) was added for 6–12 h immediately preceding the start of an experiment.

Isolation and Activation of Mouse Lymphocytes—Peripheral lymph nodes were isolated from BALB/c-FoxP3eGFP reporter mice (60). T cells were enriched using the Pan T cell isolation kit and AutoMACS magnetic bead separation (Miltenyi BioTech, San Diego, CA). Recovered T cells were stained for CD4-PE (clone GK1.5, BD Pharmingen) and purified by fluorescence activated cell sorting (FACS) to isolate CD4⁺FoxP3eGFP⁺ and CD4⁺FoxP3eGFP⁻ populations. To activate cells, 1.5 × 10⁵ CD4⁺FoxP3eGFP⁺ and 2.5 × 10⁵ CD4⁺FoxP3eGFP⁻ purified T cells were cultured in complete RPMI medium (2 mM glutamine, 1:100 non-essential amino acids, 1 mM sodium pyruvate, 10 mM HEPES, 100 units/ml of penicillin and streptomycin, 50 μM 2-mercaptoethanol, final concentrations) supplemented with 10% FBS and 100 units/ml of recombinant human interleukin-2 (rhIL-2). T cells were activated using plate-bound αCD3 (1 μg/ml) and αCD28 (2 μg/ml). Pf% was measured 24–72 h post-activation.

Culture and Transfection of HEK293 Cells—Standard methods of transfection and cell culture were used to express genes in HEK293 cells, as previously described (61, 62). HEK293 cells were maintained in exponential growth in Dulbecco's modified Eagle's medium supplemented with 10% heat-inactivated FBS, 2 mM glutamine, 50 units/ml of penicillin G, and 50 μg/ml of streptomycin, and incubated at 37 °C in a humidified atmosphere with 5% CO₂. The cells were enzymatically dissociated upon reaching 70–80% confluence in 75-cm² tissue culture flasks, and co-transfected with a P2X7R plasmid (2 μg) and a fluorescent reporter gene (0.5 μg) using Effectene (Qiagen, Valencia, CA). They were then replated in 35-mm culture dishes, and left to grow for 48 h before use in experiments. A 1:3 ratio of genes encoding the human P2X7aR (2 μg) and P2X7bR (6 μg) splice variants were used in co-expression experiments.

Imaging Ca²⁺ in a GCaMP5G Cell Line—HEK293 cells stably expressing GCaMP5G and transiently expressing AsRed were plated on glass coverslips for investigation using a fluorescent imaging system. GCaMP5G fluorescence was imaged using a CoolSnap EZ camera (Photometrics, Tucson, AZ) and the appropriate filter cube (excitation 480 nm; emission 510 nm). Imaging data were analyzed using μManager 1.4 (63).

Measuring Pf%Detailed descriptions of our method are described elsewhere (1, 62). Briefly, agonist-gated current (black trace in Fig. 1B) was measured electrically and then integrated (red trace in Fig. 1B) to give the total charge transfer across the cell membrane (Q_T, in coulombs). At the same time, Ca²⁺ influx was monitored from the fluorescence emitted by a high concentration (2 mM) of intracellular fura-2 K_s (gray trace

in Fig. 1B) introduced through the recording electrode. When all of the Ca²⁺ entering the cell was captured by the dye, then the change in 510 nm emission of fura-2 excited by 380 nm light (ΔF₃₈₀) was proportional to the Ca²⁺ flux through the pore (64). The ΔF₃₈₀ signal was measure in bead units and calibrated in separate experiments as previously described (1), allowing ΔF₃₈₀ to be converted to coulombs of Ca²⁺ charge (Q_{Ca}). Then, the fraction of total membrane current carried by Ca²⁺ (Pf) equaled Q_{Ca}/Q_T (slope of the line in Fig. 1C), and the percent fractional Ca²⁺ current (i.e. the Pf%) equaled P_f × 100%. In most cases, Q_{Ca} was a linear function of Q_T for the enter length of the ATP response, demonstrating that all of the incoming Ca²⁺ was captured by fura-2. Data showing non-linearity were discarded. The intracellular solution contained (in mM): 140 CsCl, 10 TEA-Cl, 10 HEPES, pH 7.3, with CsOH. The extracellular solution contained: 150 NaCl, 2 CaCl₂, 1 MgCl₂, 10 glucose, 10 HEPES, pH 7.4, with NaOH. MgCl₂ was omitted from the extracellular solution when 600 μM BzATP was used as agonist. Often, we were able to record several currents (2–10) from a single cell. In these cases, successive applications were separated by >2 min to minimize current facilitation (65) and the time-dependent change in deactivation (66). We saw no use-dependent change in Pf%, and therefore multiple determinations were averaged to yield a single value.

As noted above, the Pf% method requires that all the Ca²⁺ that enters the cell is captured by the high concentration of intracellular fura-2 (49). In most cases, this condition is satisfied using short applications of submaximal concentrations of agonist (1). The rat P2X7kR is an exception because its unusually slow deactivation produces sustained inward currents and Ca²⁺ fluxes (67, 68) that eventually saturate the intracellular fura-2. We found that the fura-2 remained unsaturated during the first 5–20 s of inward current because Ca²⁺ flux through the P2X7kR was unusually small (see below). Thus, all Pf% measurements of rat P2X7kRs are calculated from the initial, unsaturated portion of the response occurring in the first 20 s or less of the BzATP-gated inward current.

Calculating P_{Ca}/P_{Na}—We used our empirical measurements of Pf% to estimate P_{Ca}/P_{Na} using the following equation (54),

$$\frac{P_{Ca}}{P_{Na}} = \frac{\gamma_{Na} \times [Na]_o - \frac{P_{Cs}}{P_{Na}} \times \gamma_{Cs} \times [Cs]_i \times \exp\left(\frac{2VF}{RT}\right)}{4 \times \gamma_{Ca} \times [Ca]_o \times \left(\frac{1}{Pf} - 1\right)} \quad (\text{Eq. 1})$$

where γ_{Na}, γ_{Cs}, and γ_{Ca} are the activity coefficients for Na⁺ (0.72), Cs⁺ (0.72), and Ca²⁺ (0.57), respectively. The concentrations of extracellular Na⁺ and intracellular Cs⁺ were those used in the Pf% experiments (see above), and the concentration of extracellular free Ca²⁺ was calculated using MaxChelator (69). P_{Cs}/P_{Na} equaled 1 (70). F, R, and T had their usual values at 22 °C, and V equaled –60 mV.

Measuring P_{NMDG}/P_{Na}—We used a standard reversal potential approach, described in detail in Migita *et al.* (71), to measure relative N-methyl-D-glucamine⁺ (NMDG⁺) permeability (i.e. P_{NMDG}/P_{Na}). Voltage ramps (140 mV, 200 ms) were applied

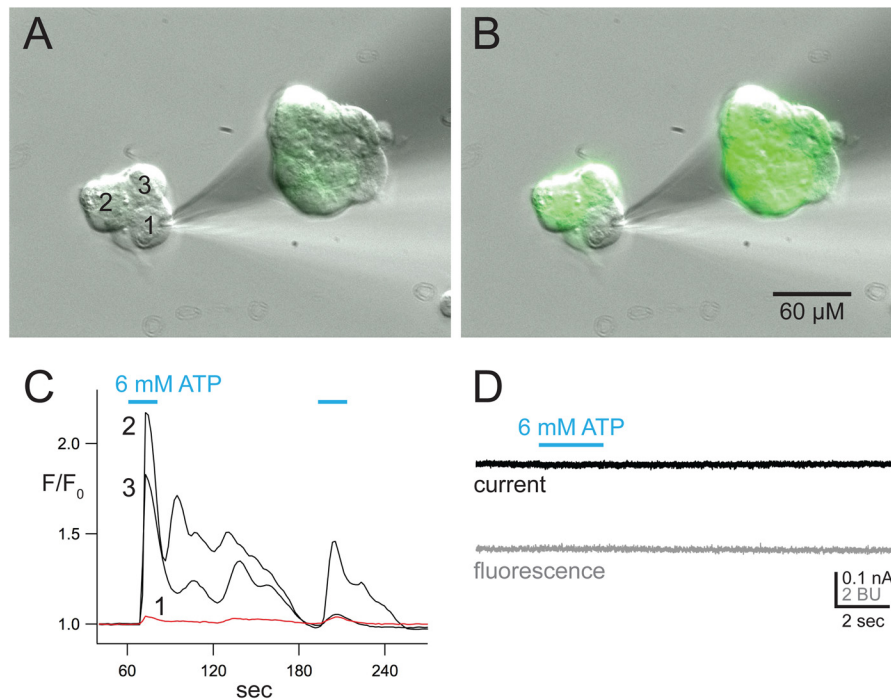


FIGURE 2. **The metabotropic P2Y response is missing in dialyzed cells.** *A*, resting HEK293-GCaMP5G cells show low levels of fluorescence. Cell 1 is undergoing internal dialysis with the contents of a recording electrode used for whole cell, broken-patch electrophysiology. The electrode contained the normal intracellular fura-2 solution used to measure *Pf*%. *B*, fluorescence increases dramatically in Cells 2 and 3 in response to activation of endogenous P2Y receptors by bath application of 6 mM ATP. The change in fluorescence intensity indicates an increase in [Ca²⁺]_i. In contrast, Cell 1 shows no significant change in fluorescence. *C*, the change of fluorescence intensity was measured from each the three cells in the cluster. The image capture was started 5 min after the start of intracellular recording, and proceeded at a rate of one capture per 2 s. The change in fluorescence intensity was quantified by dividing the intensity of each capture by the average resting intensity measured before drug application. ATP was applied twice (cyan bars). *D*, current (top) and fura-2 *F*₃₈₀ fluorescence (bottom) was captured from a single naive HEK293 using our standard method of measuring *Pf*%. ATP caused no measurable change in *F*₃₈₀. BU, bead units.

to HEK293 cells expressing P2X7Rs before and during short applications of BzATP (10–100 μM). The holding voltage was –100 mV, and the extracellular solution contained (in mM): 150 mM XCl, 2 CaCl₂, 1 MgCl₂, 10 HEPES, and 10 glucose, pH 7.4, where *X* was either Na⁺ or NMDG⁺. Calcium and magnesium were included to replicate the conditions used to measure *Pf*%. The intracellular solution contained (in mM): 150 NaCl, 10 EGTA, 10 HEPES, pH 7.3. Relative NMDG⁺ permeability (*P*_{NMDG}/*P*_{Na}) was determined from the difference in reversal potentials of agonist-gated currents measured in the two extracellular solutions ($\Delta E_{\text{rev}} = E_{\text{rev,NMDG}} - E_{\text{rev,Na}}$) as,

$$\frac{P_{\text{NMDG}}}{P_{\text{Na}}} = \frac{[\text{Na}]_i \times \gamma_{\text{Na}} \times \exp\left(\frac{\Delta E_{\text{rev}} F}{RT}\right) - 4 \times [\text{Ca}]_o \times \gamma_{\text{Ca}} \times \frac{P_{\text{Ca}}}{P_{\text{Na}}}}{[\text{NMDG}]_o \times \gamma_{\text{NMDG}}} \quad (\text{Eq. 2})$$

where γ_{NMDG} equaled 0.72, and *P*_{Ca}/*P*_{Na} was calculated from the *Pf*% as described above (see Equation 1). ΔE_{rev} is the difference in reversal potentials measured in extracellular solutions containing either Na⁺ and Ca²⁺ or NMDG⁺ and Ca²⁺.

Homology Modeling of Resting and Activated rP2X7aRs—Models for the rat P2X7 receptor were built as previously described (72). Briefly, the apo (Protein Data Bank 4DW0) and ATP-bound (Protein Data Bank 4DW1) structures of zP2X4.1 (73) were used as templates to generate the rP2X7aR models using the homology modeling server SWISSMODEL (74). The rP2X7aR models were minimized using the GROMOS force-

field implemented in the program DEEP VIEW (75). Models were visualized in PyMol.

Data Analysis—We used GraphPad Prism for statistical analyses. The averaged values of pooled data with normal distributions are shown as mean ± S.E. Unless otherwise noted, groups were analyzed by one-way analysis of variance with significance determined by Tukey's multiple comparison test. The exceptions were analyzed using Student's *t* test. *p* values of ≤ 0.01 were considered statistically significant.

RESULTS

HEK293 cells express metabotropic P2Y receptors capable of increasing the concentration of intracellular Ca²⁺ ([Ca²⁺]_i) by mobilizing internal Ca²⁺ stores in a GTP-dependent manner (76, 77). We used a HEK293 cell line with stable expression of the genetically encoded Ca²⁺ sensor, GCaMP5G (78), to determine whether activation of P2Y receptors by ATP (6 mM) is an unintended consequence of using high agonist concentrations to measure *Pf*%. We found that ATP increases [Ca²⁺]_i in unperturbed cells, but failed to do so in cells whose cytoplasm is dialyzed with the contents of a whole cell recording electrode (Fig. 2, A–C). Furthermore, when using our standard method of measuring *Pf*%, we saw no change in fura-2 fluorescence in mock transfected cells (*n* = 11) in response to a concentration of agonist (6 mM) that was 10–60 times higher than that applied to functional recombinant P2X7Rs in our study (Fig. 2D). These data support the contention that the internal dialysis that occurs during whole cell *Pf*% recordings disrupts the

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TABLE 1

Measured and adjusted Pf% values

Pf% was measured from HEK293 cells expressing the specific P2X7R listed in the leftmost column. Concentrations are mM. Species are homomeric “a” splice variants unless otherwise noted in parentheses. 0.1 mM BzATP was dissolved in an extracellular solution containing 2 mM CaCl₂ and 1 mM MgCl₂, 0.6 mM BzATP was dissolved in an extracellular solution containing 2 mM CaCl₂ and no added MgCl₂. The estimated concentrations of free Ca²⁺ and Mg²⁺ were calculated using MaxChelator. Pf%_{adj} is the fractional Ca²⁺ current expected if Pf% could be measured in the absence of Ca²⁺ chelation by the purinergic agonist. P_{Ca}/P_{Na} was calculated from the empirically measured Pf% values as described under “Experimental Procedures.”

Species	[BzATP]	[Ca ²⁺] _{free}	[Mg ²⁺] _{free}	Pf%	Pf% _{adj}	P _{Ca} /P _{Na}	n
Dog	0.1	1.95	0.95	7.3 ± 0.2	7.5 ± 0.2	1.8 ± 0.0	11
Rat	0.1	1.95	0.95	7.1 ± 0.2	7.3 ± 0.2	1.7 ± 0.1	136
Rat (k)	0.1	1.95	0.95	2.8 ± 0.3 ^a	2.8 ± 0.3 ^a	0.7 ± 0.1 ^a	34
Mouse	0.1	1.95	0.95	7.1 ± 0.4	7.3 ± 0.4	1.7 ± 0.1	11
Mouse (k)	0.1	1.95	0.95	4.3 ± 0.3 ^b	4.4 ± 0.3 ^b	1.0 ± 0.1 ^b	13
Guinea pig	0.6	1.40	0	5.7 ± 0.8	8.2 ± 1.1	1.9 ± 0.3	7
Monkey	0.6	1.40	0	8.0 ± 0.4	11.4 ± 1.0	2.7 ± 0.3	8
Human	0.6	1.40	0	7.1 ± 0.4	10.2 ± 0.6	2.4 ± 0.2	22
Human X7 (a/b)	0.6	1.40	0	8.2 ± 0.3	11.7 ± 0.4	2.9 ± 0.1	8
Zebrafish	0.6	1.40	0	14.3 ± 1.0 ^c	20.4 ± 1.4 ^c	5.6 ± 0.5 ^c	14

^a Significantly different from rP2X7aR.

^b Significantly different from mP2X7aR.

^c Significantly different from all other species.

P2Y response (1). Confident that we could record a pure ionotropic response, we then moved to the study of the P2X7Rs.

Genes encoding the P2X7aRs of dog, frog, guinea pig, human, monkey, mouse, rat, and zebrafish were individually expressed in HEK293 cells for study using patch-clamp photometry. Cells expressing transfected genes were visually identified by the fluorescence emitted from a reporter protein (eGFP or dsRed), and the Pf% was determined as described above. Transfected cells responded with robust inward currents (membrane holding potential = -60 mV) and changes in fura-2 fluorescence to applications of BzATP (100–600 μM) with one exception: we failed to record agonist-gated current or a change in fura-2 fluorescence in response to BzATP (up to and including 1 mM) and ATP (6 mM) in HEK293 cells transfected with the gene encoding the *Xenopus laevis* P2X7aR. This was surprising because injection of frog P2X7aR cRNA results in large ATP-gated currents in a *Xenopus* oocyte expression system (79).

Most P2X7aRs Transduce Substantial Ca²⁺ Currents—In all other cases, we reliably measured the Pf% with the dye-overload technique. First, we tested the P2X7aR constructs using a concentration of BzATP (100 μM) that does not significantly chelate Ca²⁺ (estimated free [Ca²⁺]_o equaled 1.95 mM) (8). Only three (mouse, rat, and dog) of the seven responded with sizable membrane currents and changes in fura-2 fluorescence. In each case, the average Pf% equaled ~7% (Table 1). For the mouse ortholog, we studied both the Pro⁴⁵¹ and Leu⁴⁵¹ alleles (80, 81), and found no significant difference in their Pf% values.

Next, we used a higher concentration of BzATP (600 μM) and a Mg²⁺-free test solution (9, 48) to elicit responses from the remaining four constructs (zebrafish, guinea pig, monkey, and human). Under these conditions, the calculated free [Ca²⁺]_o drops from 1.95 to 1.40 mM as the [BzATP] rises from 0.1 to 0.6 mM and the [Mg²⁺]_o falls from 1.0 to 0.0 mM. Although less Ca²⁺ was available to carry the agonist-gated current, we still measured substantial Pf% values equal to ~6–8% for guinea pig, monkey, and human P2X7aRs (Table 1). The Pf% of the zebrafish P2X7R was higher still at 14.3 ± 1.0% (n = 14).

Pf% Adjusted for Changes in [Ca²⁺]_o—We sought to determine a fair method of comparing Pf% values measured under conditions where the free [Ca²⁺]_o varied as a consequence of the agonist concentration. To this end, we measured the Pf%

values of rat P2X7aR currents elicited by 100 μM BzATP and a range of ATP concentrations (Fig. 3, A and B). We used ATP because the cost of using millimolar concentrations of BzATP was prohibitive, and because others have shown that ATP and BzATP bind Ca²⁺ equally well (8). Furthermore, ATP is the endogenous agonist for native receptors, and thus provides insight regarding the amplitude of the Ca²⁺ flux under pathophysiological conditions where the high [ATP]_o causes significant buffering of extracellular Ca²⁺. We calculated the extracellular concentration of free Ca²⁺ in a given concentration of BzATP or ATP using MaxChelator (Fig. 3C), and then empirically determined the Pf%. As expected, the Pf% progressively decreased with increasing concentrations of agonist (Fig. 3D), falling from a high of 7.1 ± 0.2% (n = 136) at 0.1 μM BzATP to a low of 1.7 ± 0.3% (n = 12) at 6 mM ATP. The XY-plot of Pf% versus free [Ca²⁺]_o was linear (Pearson's r = 0.99) with a slope of 3.8% per mM extracellular Ca²⁺ (Fig. 3E).

We took advantage of this linear relationship to calculate an adjusted Pf% (Pf%_{adj}) equal to the product of Pf% and R, where R is the ratio of the free [Ca²⁺]_o values expected in the absence (2 mM) and presence of 100 or 600 μM BzATP. The Pf%_{adj} values are ~3 and 43% higher than the empirically measured Pf% values for 100 and 600 μM BzATP, respectively (Table 1), and thus represent the projected contribution of Ca²⁺ to the total agonist-gated cation current measured in the absence of Ca²⁺ chelation (Fig. 3F). We used a multiple comparisons test to detect differences among the mean Pf%_{adj} values of all seven species and found that one species differed from all others: that is, Ca²⁺ makes a greater adjusted contribution (~20%) to the BzATP-gated current of the zebrafish P2X7aR than it does in any other species. Indeed, the unadjusted, empirically determined Pf% measured in the relatively low [Ca²⁺]_o of 1.4 mM is still larger than that of other members of the ligand-gated ion channel superfamily measured in a higher [Ca²⁺]_o typically equal to 1.8 mM (82).

We investigated the cause of the transcendent Pf% of the zebrafish P2X7aR using a site-directed mutagenesis approach. The zebrafish P2X7aR retains the polar amino acids of TM2 that facilitate Ca²⁺ flux through rat P2X2Rs (1, 71), but lacks the vestibular acidic amino acids that are responsible for the high Pf% values of rat and human P2X1Rs and P2X4Rs (83, 84).

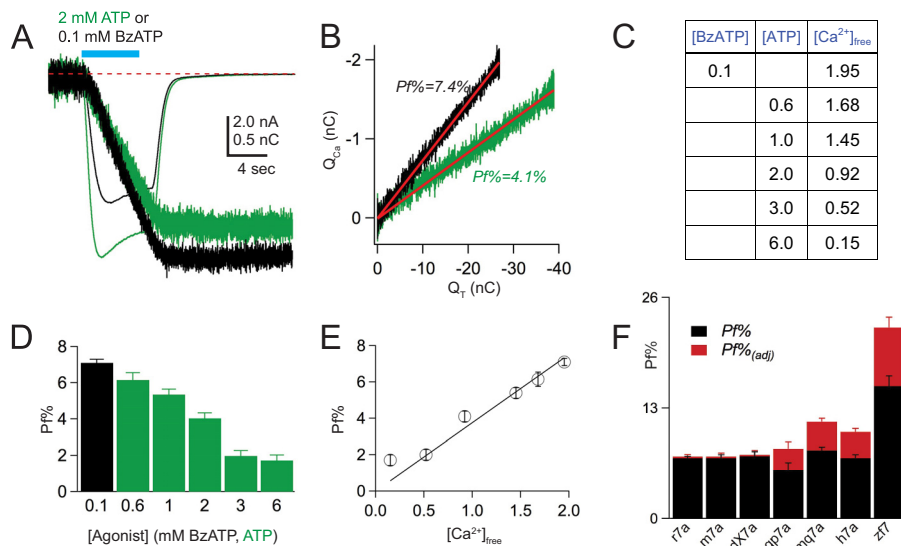


FIGURE 3. **Pf% decreases with increasing concentrations of ATP.** A, 2 mM ATP (green traces) evokes a larger current and smaller Q_{Ca} (shown in nC after conversion from bead units; see "Experimental Procedures") in an HEK293 cell expressing the rat P2X7aR than does 100 μ M BzATP (black traces). B, the Pf%, determined from the linear fit of Q_{Ca} versus Q_T , is smaller for currents evoked by 2 mM ATP by comparison to 100 μ M BzATP, despite the fact that the currents are larger. C, the calculated free $[Ca^{2+}]_o$ in different concentrations of BzATP and ATP. In these calculations, we used concentrations of total Na^+ , Ca^{2+} , and Mg^{2+} equal to 150, 2, and 1 mM, respectively, a pH of 7.4, and a temperature of 23 °C. D, the histogram shows the average values \pm S.E. of the Pf% of the rat P2X7aR measured using 100 μ M BzATP (black bar) and increasing concentrations of ATP (green bars). The Pf% decreases with increasing concentrations of ATP. E, the graph plots empirical Pf% against the calculated free $[Ca^{2+}]_o$. The result is a straight line, suggesting that the decrease in Pf% reflects chelation of extracellular Ca^{2+} by the purinergic agonists. F, the histogram shows the Pf% (black bars) and Pf%_{adj} (red bars) for a range of P2X7Rs.

We considered the possibility that two glutamates (Glu⁴⁸ and Glu⁵¹) just downstream of the zebrafish TM1 might take the place of the vestibular acidic amino acids of rat and human P2X1R and P2X4R that are missing in the P2X7Rs, and thus provide an electrostatic potential capable of concentrating Ca^{2+} at the entrance to the pore. To test this hypothesis, we removed the negative charge by mutating both glutamates to glutamine. Although the resulting mutant (zfP2X7aR-E48N/E51N) showed currents in response to 600 μ M BzATP that were comparable in size to that of the wild-type zfP2X7R, the concomitant change in fluorescence was reduced. From these data, we calculated a Pf% equal to $8.0 \pm 0.5\%$ ($n = 8$), which is significantly smaller than that of the wild-type zebrafish receptor ($\sim 14\%$) and more in line with those of the other P2X7aR orthologs. This result suggests that acidic amino acids play a role in regulating the Ca^{2+} flux of the zfP2X7aR, perhaps by pulling Ca^{2+} into the lateral portals that form the entrance to the pore (72).

Co-transfection of hP2X7aR and hP2X7bR—Heteromeric hP2X7a/bRs show channel properties in fluorescent assays that differ from those of the homomeric hP2X7aR (56). To determine whether heteromerization affects Ca^{2+} transport, we measured the Pf% of the BzATP-gated current of HEK293 cells transfected with genes encoding P2X7aRs and hP2X7bRs. Cells expressing only the P2X7bR showed small membrane currents (< 2 pA/pF) in response to relatively high concentrations of BzATP (600 μ M) and ATP (6 mM). The fluorescent signals associated with these currents were too small to quantify. In contrast, 600 μ M BzATP evoked large currents (> 50 pA/pF) in cells co-transfected with a 1:3 ratio of the hP2X7A and hP2X7B genes. These currents resembled in size and shape those recorded from cells expressing only the hP2X7aR. We compared the ATP concentration-current curves of cells trans-

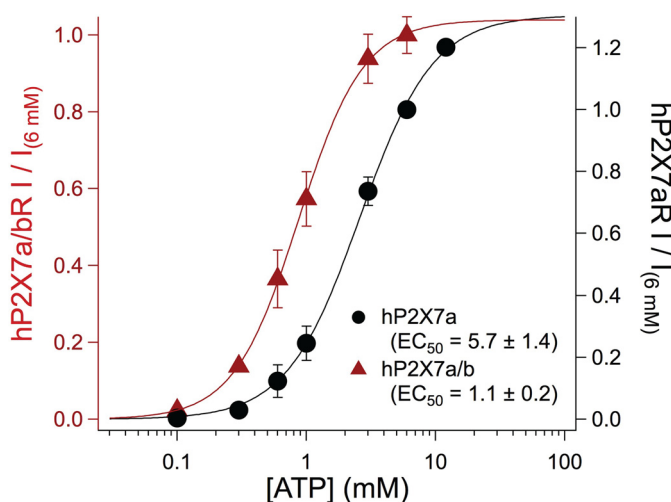


FIGURE 4. **Concentration-current curves for homomeric and heteromeric human P2X7Rs.** Concentration-response curves were built from HEK293 transfected either with the hP2X7aR plasmid alone (black), or co-transfected with the hP2X7aR and hP2X7bR plasmids (red). Each data point was normalized to the peak current caused by applying 6 mM ATP to the same cell. Individual data points corresponding to specific concentrations of ATP were averaged (solid circles). The solid lines show the best fit of the averaged data to the Hill equation. For statistical analysis, EC_{50} values were calculated from data of complete concentration-response curves obtained from 12 individual cells. Significance was determined using Student's t test.

ected with one or both genes to gauge heteromerization using a low-divalent (0.2 Ca^{2+} and no Mg^{2+}) extracellular solution (8, 9). In keeping with Adinolfi *et al.* (56), we saw leftward shifts in the concentration-current curves of cells transfected with both genes by comparison to cells expressing only the P2X7aR (Fig. 4). From these data, we measured a significant difference (Student's t test; $p < 0.01$) in agonist potency of cells transfected with one (hP2X7A; ATP $EC_{50} = 5.3 \pm 1.4$ mM, $n = 6$) or both

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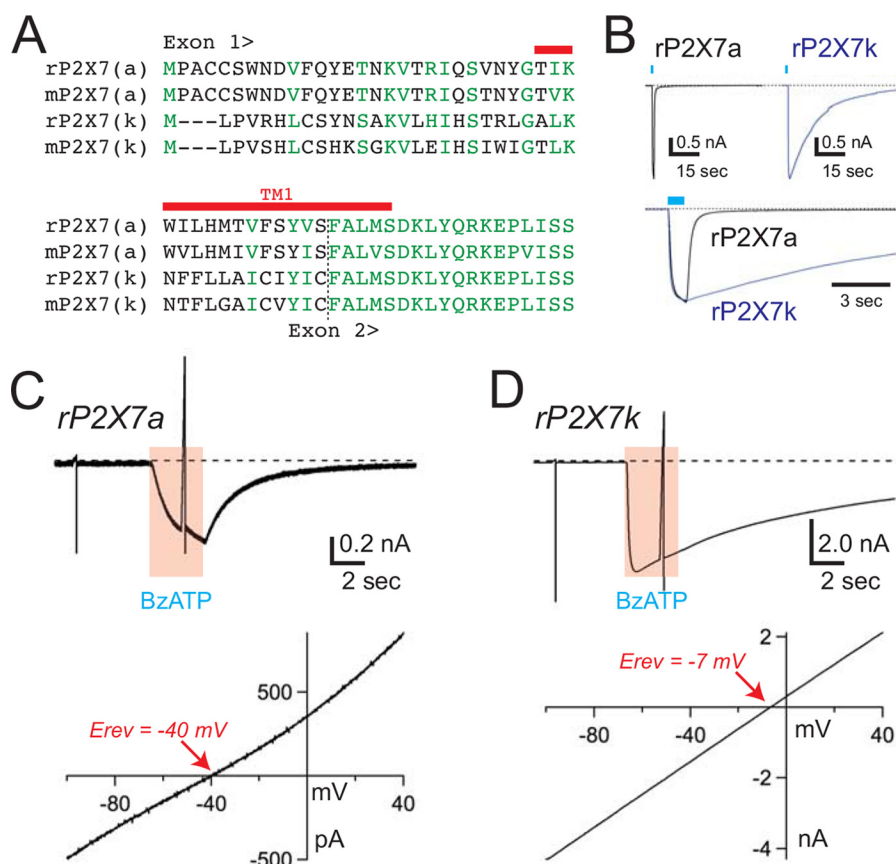


FIGURE 5. The rP2X7kR is constitutively dilated. *A*, sequence alignment of the N terminus and TM1 of rat and mouse P2X7a and P2X7k receptors. Identical and similar residues are shown in *green*. TM1 is marked with a *red bar*. *B*, the *top two traces* show the time courses of the membrane currents gated by 100 μ M BzATP in voltage-clamped HEK293 cells (holding voltage = -60 mV) expressing either rP2X7aRs (*left*) or rP2X7kRs (*right*). The rP2X7kR often took several minutes to fully deactivate upon washout of agonist. In the *bottom trace*, normalized rP2X7aR and rP2X7kR currents are overlaid to show the difference in deactivation times. *C* and *D*, voltage ramps (140 mV, 200 ms) from a holding voltage of -100 mV were applied to HEK293 cells expressing either rP2X7aRs (*panel C*) or rP2X7kRs (*panel D*) before and during applications of 30 μ M BzATP. Currents through the rP2X7aR were reversed at membrane potentials (*bottom left trace*) that were significantly more negative than the currents through the rP2X7kR (*bottom right trace*). These data support the hypothesis that the pore of the rP2X7kR is constitutively dilated. Similar results were found using the mouse splice variants.

(*hP2X7A/hP2X7B*; ATP EC₅₀ = 1.1 \pm 0.2 mM, *n* = 6) genes, suggesting that some of the agonist-gated current flows through a population of heteromeric receptors. We then measured the *Pf*% of cells co-transfected with genes encoding hP2X7aRs and hP2X7bRs. Although the *Pf*% values measured from these cells tended to be higher (8.2 \pm 0.5%, *n* = 8) than cells expressing P2X7aR alone (\sim 7.1%; see Table 1), the difference in the two values failed to reach statistical significance (Student's *t* test; *p* = 0.1229). Although many factors can influence the outcome of co-expression experiments, our results suggest that heteromerization affects the binding/gating properties of human P2X7Rs (*i.e.* the EC₅₀) more than ionic selectivity (*i.e.* the *Pf*%).

The Murine P2X7k Splice Variant Shows Limited Ca²⁺ Flux—Rat and mouse P2X7kRs are derived from an alternative exon 1' (6) and contain unique N termini and TM1s that differ from those of the P2X7aR splice variants (Fig. 5*A*). We determined the *Pf*% values of the P2X7kR splice variants in response to the same concentration of BzATP (100 μ M) used to study the P2X7aRs. We studied the rat P2X7kR first, and in keeping with previous reports (57), we found two obvious differences in the character of its BzATP-gated currents. First, deactivation of the transmembrane current was noticeably slower than that of

the rat P2X7aR (Fig. 5*B*). Second, the reversal potential of the BzATP-gated current measured in an extracellular solution containing 150 mM NMDG⁺, 2 mM Ca²⁺, and 1 mM Mg²⁺ was right-shifted by \sim 25 mV (-41.8 ± 0.6 mV for the rP2X7aR, and -15.0 ± 2.4 mV for the rat P2X7kR; Fig. 5, *C* and *D*). These data yield $P_{\text{NMDG}}/P_{\text{Na}}$ of 0.12 \pm 0.00 (*n* = 12) and 0.55 \pm 0.05 (*n* = 8) for the rat P2X7aR and P2X7kR, respectively. A similar difference was found for the mouse P2X7aR and P2X7kR ($P_{\text{NMDG}}/P_{\text{Na}}$ 0.05 \pm 0.01, *n* = 11; and 0.40 \pm 0.03, *n* = 11). The disparity in baseline permeability to the large (4.5 Å) polyatomic NMDG⁺ supports the suggestion that the selectivity filters of the P2X7a and P2X7k splice variants adopt different initial open state conformations with dissimilar cation permeabilities (57). To explore this hypothesis further, we sought to determine whether the contribution of the physiologically relevant divalent cation, Ca²⁺, to whole cell current also differed.

We measured the *Pf*% of the BzATP-gated current of the rat and mouse P2X7kRs. In both instances, 100 μ M BzATP evoked inward membrane currents and changes in fura-2 fluorescence emission as expected for non-selective cation channels permeable to Ca²⁺ (for example, see Fig. 6*A*). Within the constraints outlined under "Experimental Procedures," we measured a linear relationship of Q_{Ca} versus Q_{T} (Fig. 6*B*), as expected if all of

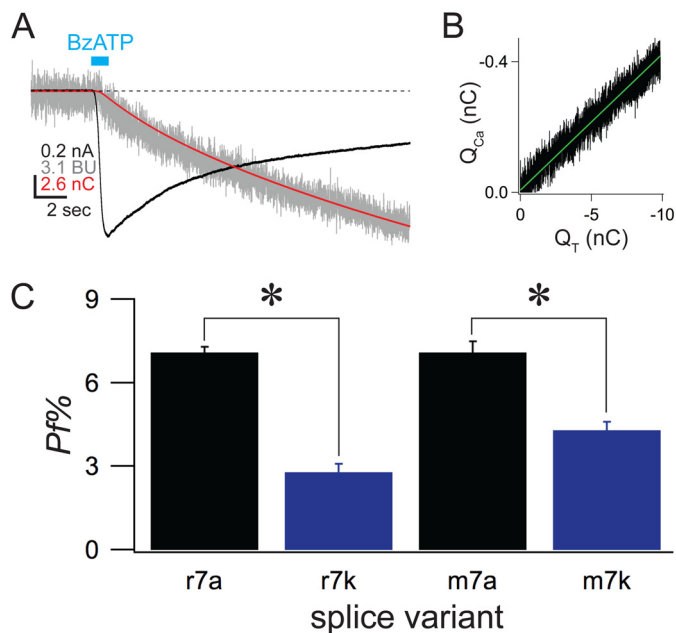


FIGURE 6. The rP2X7kR carries a small Ca²⁺ current. *A*, the traces show raw data of an experiment that measured the *Pf*% of a cell expressing the rP2X7kR. The cell was held at -60 mV under voltage-clamp. BzATP ($100 \mu\text{M}$) was applied for the time indicated by the cyan bar. The black trace is the membrane current measured in nA. The red trace is the integrated membrane current and equals Q_T . The gray trace is the normalized fura-2 fluorescence, measured in bead units and proportional to Q_{Ca} (see "Experimental Procedures" for full details). *B*, the graph shows a linear relationship between Q_T (x axis) and fluorescence (y axis), a strict requirement of the dye-overload method (64). *C*, the histogram shows the average data of mouse and rat splice variants.

the Ca²⁺ that binds to the intracellular fura-2 comes from the ATP-gated inward current traveling through the channel pore. We discovered that the *Pf*% of the agonist-gated inward current of the rat P2X7kR ($2.8 \pm 0.3\%$; $n = 34$) was significantly smaller than that of the rat P2X7aR (7.1%). Again, similar results were found when comparing splice variants from mouse (4.3 versus 7.1% for the P2X7kR and P2X7aR splice variants, respectively; see Fig. 6C and Table 1). Taken together, our measurements of $P_{\text{NMDG}}/P_{\text{Na}}$ and *Pf*% show that P2X7kRs conduct smaller Ca²⁺ currents than rP2X7aRs despite having a higher permeability to large cations, implying that a critical Ca²⁺ selectivity filter is disrupted in the constitutively dilated P2X7kRs. These findings bring forth two intriguing hypotheses. First, P2X7a and P2X7k splice variants differ only in the composition of their N termini and TM1s. Their pore forming TM2 domains are identical, suggesting that they cannot be the primary sites responsible for the different *Pf*% of the two splice variants. Thus, our results suggest that domains that lie outside of the ion-conducting pathway of the P2X7R are capable of selectively regulating its Ca²⁺ current. Second, the physiologically relevant outcome of pore dilation may not be an increased permeability to large cations as generally assumed, but rather a decreased permeability to Ca²⁺, an essential trigger of the downstream sequela of P2X receptor activation (5).

Relative Ca²⁺ Permeability—Relative Ca²⁺ permeabilities are commonly used to compare the selectivity of ligand-gated ion channels for Ca²⁺ (82, 85). Therefore, we used our empirical measurements of *Pf*% to estimate the relative Ca²⁺ to Na⁺ permeability (P_{Ca}/P_{Na}) within the limits of the Goldman-Hodgkin-

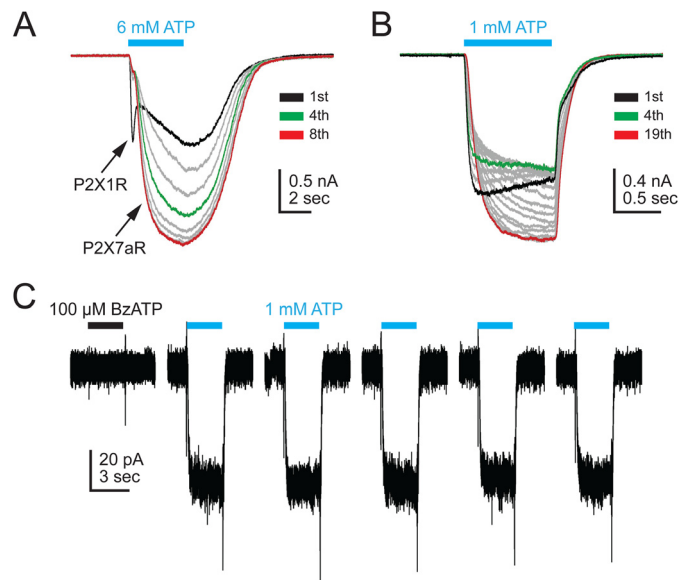


FIGURE 7. Native currents of immune cells. *A*, human macrophage. ATP was applied every 60 s for the time indicated by the solid cyan bar. The first application (black trace) shows a mixed response with P2X1R and P2X7aR components. Subsequent traces showed predominately P2X7aR currents. The P2X1R response was blocked by an antagonist during *Pf*% measurements (see "Results"). The P2X7aR response showed current facilitation similar to that described for many recombinant P2X7aRs. Green and red traces equal the fourth and eighth applications of ATP, respectively. *B*, mouse macrophage. The native response also showed facilitation. *C*, mouse regulatory T lymphocyte. $100 \mu\text{M}$ BzATP (black bar) had no effect on membrane current. In contrast, repetitive applications of ATP (1 mM) evoked steady currents that showed no facilitation.

Katz equation (64), allowing the calcium selectivity of P2X7Rs to be compared with those of other P2XR subtypes and other classes of ligand-gated ion channels. The derived P_{Ca}/P_{Na} of the P2X7Rs equaled 0.7 to 5.6 (Table 1), values that approximate those of the empirically measured P_{Ca}/P_{Na} of other P2XR subtypes (1.2–4.8) and other ligand-gated channels (0.4–10.4) (82, 86, 87). As expected, the upper and lower limits of the purinergic P_{Ca}/P_{Na} were set by the zfpP2X7aR and the murine P2X7kRs, respectively.

The *Pf*% of Native P2X7 Channels of Mouse and Human Immune Cells—We determined the *Pf*% of the BzATP-gated current of monocyte-derived macrophages isolated from mouse peritoneum and human blood (Fig. 7, *A* and *B*). After a minimum of 6 days in culture, we incubated the macrophages with $1 \mu\text{g}/\text{ml}$ of lipopolysaccharide for 6–12 h and then measured their response to 100 (mouse) or $600 \mu\text{M}$ (human) BzATP. The purinergic antagonist, 2',3'-*O*-(2,4,6-trinitrophenyl)-ATP (100 nM), was included in the superfusate at the time of the experiment to block the possible contribution of P2X1Rs to the BzATP-gated current (see Fig. 7*A*) (88). Measured in this way, the *Pf*% of the ATP-gated current of mouse and human macrophages equaled 8.8 ± 1.0 ($n = 7$) and 9.0 ± 0.5 ($n = 10$), respectively. We then used the empirical measurements to calculate $Pf\%_{\text{adj}}$ and P_{Ca}/P_{Na} values. For mouse macrophages, $Pf\%_{\text{adj}}$ equaled $9.0 \pm 1.0\%$ and P_{Ca}/P_{Na} equaled $2.3 \pm 0.3\%$. For human macrophages, $Pf\%_{\text{adj}}$ and P_{Ca}/P_{Na} equaled 12.8 ± 0.8 and $3.3 \pm 0.2\%$, respectively. Macrophages preferentially express the P2X7aR (68). In keeping with this observation, we found no statistically significant difference in the *Pf*% measured from recombinant P2X7aRs expressed in HEK293 cells and

The Ca²⁺ Current of P2X7 Receptors

those measured from the native ATP-gated responses of species-matched macrophages. To the best of our knowledge, these values are the first quantitative measures of the Ca²⁺ current of a native P2X7aR, and prove the hypothesis that P2X7Rs transduce significant Ca²⁺ currents in immune cells.

Finally, we measured the *Pf*% of the ATP-gated current of mouse CD4⁺Foxp3⁻ conventional and CD4⁺Foxp3⁺ regulatory T-lymphocytes that preferentially express the P2X7kR splice variant (68), using the same extracellular divalent cation concentrations (2 mM total Ca²⁺ and 1 mM total Mg²⁺) used to study recombinant P2X7kRs. Both types of resting CD4⁺ T-lymphocytes failed to respond to either 100 μM BzATP or 1 mM ATP; at these concentrations, both agonists fully activate P2X7kRs (57, 89). Likewise, activated conventional CD4⁺ T cells did not respond to either agonist with inward current. In contrast, 1 mM ATP evoked inward current (5.0 ± 1.3 pA/pF; *n* = 11) in activated CD4⁺ regulatory T cells with a *Pf*% of 5.2 ± 0.6% (*n* = 9). The adjusted *Pf*%_{adj} equaled 7.1 ± 0.6% (free [Ca²⁺]_o = 1.45; see Fig. 2C), and the calculated *P*_{Ca}/*P*_{Na} equaled 1.8 ± 0.2. These same cells failed to respond to 100 μM BzATP (Fig. 7C). The inability of BzATP to induce membrane current when applied at a concentration greatly exceeding its EC₅₀ (~8 μM; see (57)), and the unexpectedly high *Pf*%, argue against the involvement of homomeric P2X7kRs in the CD4⁺Foxp3⁺ regulatory T cell response.

DISCUSSION

The allosteric block of P2X7R current by extracellular divalent cations complicates direct *P*_{Ca}/*P*_{Na} measurements, making it difficult to judge the relative merit of this subfamily as a Ca²⁺ source by comparison to other P2XRs and the ligand-gated ion channel superfamily as a whole. We circumvented this problem by measuring the fractional contribution of Ca²⁺ to the total ATP-gated membrane current (*i.e.* the *Pf*%) using a true-to-life concentration of extracellular Ca²⁺ and an appropriate membrane potential of -60 mV (90, 91). We found that empirical measurements using recombinant receptors ranged from 3 to ~15% by comparison to the average *Pf*% of the P2X family as a whole (8.6 ± 0.8%; red dotted line of Fig. 8A). However, these comparisons are not without fault. This is because the P2X7Rs of guinea pig, monkey, human, and zebrafish required a concentration of BzATP (600 μM) that was high enough to significantly lower the concentration of free [Ca²⁺]_o. We circumvented this problem by calculating a *Pf*%_{adj} that accounts for the variations in free [Ca²⁺]_o, and found that P2X7aRs take their place beside the P2X1R and P2X4R as family members that show significant Ca²⁺ currents. In contrast, the P2X7kRs, with the *Pf*%s (and *Pf*%_{adj}) of ~3–4%, resemble those of P2X3Rs and P2X5Rs in showing no appreciable preference (calculated *P*_{Ca}/*P*_{Na} ≤ 1) for Ca²⁺ over Na⁺.

By measuring *Pf*% values, we were able to estimate *P*_{Ca}/*P*_{Na}. Our calculated values ranged from ~1 for the entirely cation non-selective P2X7kRs, to ~2–6 for the P2X7aRs that preferentially transport Ca²⁺ (see Table 1). We calculated similar ratios (~2–3) for the native responses of mouse and human macrophages and lymphocytes. All of these values are comparable with those measured from other types of ligand-gated ion channels (82), but remarkably different from that (*P*_{Ca}/*P*_{Na} =

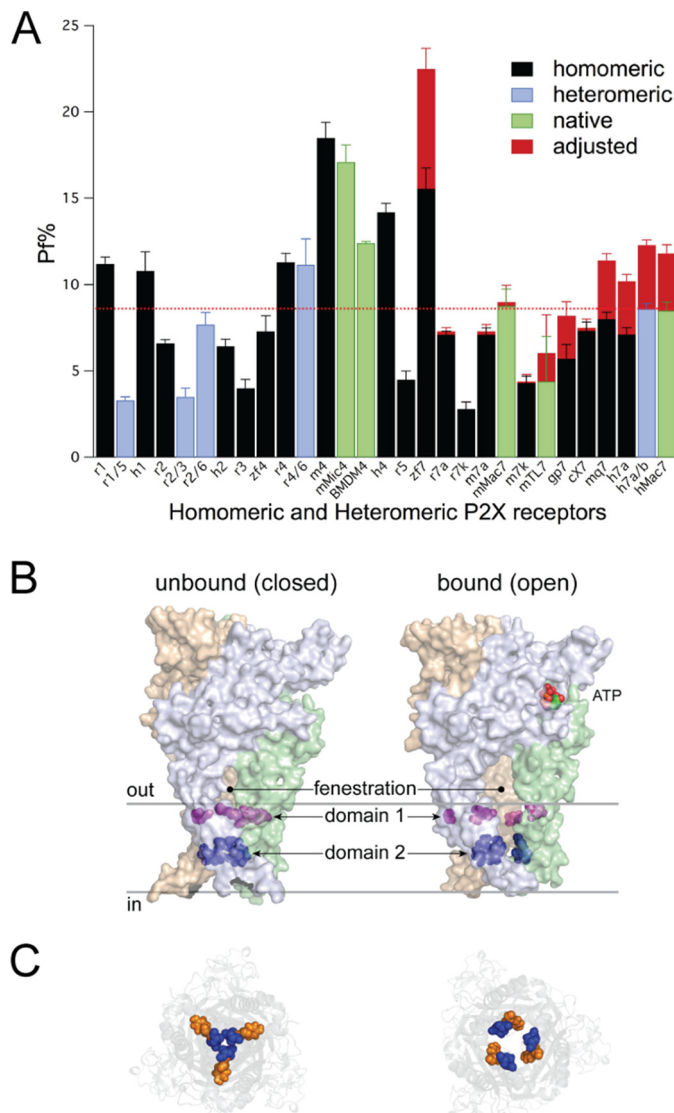


FIGURE 8. Fractional Ca²⁺ current of homo- and heteromeric P2X receptors. A, the graph shows the average *Pf*% for all homomeric (black), heteromeric (blue), and native (green) P2X receptors measured to date. Data of native and recombinant P2X7Rs come from the present study. Measures of the *Pf*% of recombinant P2X1R–P2X6Rs come from our published data (1, 83, 100) with one exception; data of recombinant hP2X2aR are previously unpublished observations. Data of the native P2X4R of mouse bone marrow-derived macrophages (BMDM4) are unpublished observations of Samways and Egan obtained during a previous study (27). The red bars indicate the average adjusted *Pf*% of the P2X7Rs expected in the absence of Ca²⁺ chelation by BzATP. The red line indicates the average *Pf*% of all P2XRs. P2X7R data come from the present study. *c*, *gp*, *h*, *m*, *mq*, *r*, and *zf* are canine, guinea pig, human, mouse, macaque, rat, and zebrafish data, respectively. *mM4* are native currents of mouse microglia. *hMac7* and *mMac7* are native currents of human and mouse macrophages, respectively. *mTL7* are native currents of mouse CD4⁺Foxp3⁺ regulatory T lymphocytes. B, homology model of the closed and open truncated rat P2X7aR in the closed and open states viewed from the plane of the membrane. The cytoplasmic N and C termini are missing. The solid gray lines indicate the inner and outer limits of the membrane. C, view from inside the cell looking out. The panel shows how Domain-2 (Ser³³⁹, Ser³⁴² (blue), and Tyr³⁴³ (orange)) form a narrow corridor that widens during opening. The distance between these residues is expected to increase further during pore dilation, which might lead to disruption of a low-affinity Ca²⁺ binding site.

35) reported in an early study (92) of the unclassified (but presumably P2X7R-mediated) ATP-gated current of transformed human B-lymphocytes. Although it is possible that this ATP-

gated current is unique, a more likely explanation is that the measurement reflects the inherent difficulties of determining relative Ca²⁺ permeability using low concentrations of extracellular Ca²⁺ in cells expressing a P2X7R susceptible to allosteric block by divalent cations. Indeed, a P_{Ca}/P_{Na} of 35 translates to a $Pf\%$ of ~60%, which is unprecedented in the ligand-gated ion channel superfamily.

Using 1 mM ATP, we measured a $Pf\%_{adj}$ of ~7% from activated mouse CD4⁺Foxp3⁺ regulatory T cells, a value that is not significantly different from that of recombinant P2X7aRs expressed in HEK293 cells (also ~7%). This is surprising because these lymphocytes contain a 30-fold excess of P2X7kR mRNA by comparison to P2X7aR (68). At present, we do not know the identity of the receptor responsible for the regulatory T cell response. Although both the relatively large $Pf\%$ and the relatively low sensitivity to ATP suggest a P2X7aR response, the inability of 100 μM BzATP to induce current in cells that previously reacted to 1 mM ATP is unusual for either a P2X7aR or P2X7kR effect. One possibility is the contribution of a heteromeric receptor of undetermined components. Future biochemical, biophysical, and pharmacological studies are needed to determine whether this hypothesis is correct.

Our finding that the splice variants of the murine P2X7Rs differ almost 2-fold in $Pf\%$ suggests that structural regulation of the Ca²⁺ current is more complicated than previously imagined. Although definitive identification of the cation selectivity filter remains a work in progress (84), we previously documented two domains that contribute to the significant Ca²⁺ permeability and current of the P2XR family (Fig. 8B). "Domain 1" is made of juxtamembrane acidic amino acids (six in total in the trimeric receptor) that are present in family members with the highest $Pf\%$ (P2X1R, P2X4R), and either absent (P2X2R, P2X5R) or shielded (P2X3R) in members with lower $Pf\%$ values (83). Although the precise mechanism of action is unknown, we proposed that Domain 1 amino acids use the electrostatic attraction of COO⁻ side chains to concentrate Ca²⁺ in the fenestrae that form the entrance to the pore (84, 93). Our present results provide additional support for this hypothesis, as we now show that the very high $Pf\%$ of the zfpP2X7R requires carboxylates at positions that are just extracellular to the transmembrane domains and expected to be in or near the mouth of the pore. Definitive proof of this hypothesis awaits the report of a high-resolution structural map of the zfpP2X7R protein.

"Domain 2" is made of three polar amino acids (nine in total) that line the narrowest part of the rat P2X2R pore (see Fig. 8B). Others and we showed that altering the hydrophobicity, volume, and/or charge of residues in Domain 2 affect cation (71) and anion permeability (94), and $Pf\%$ (1). Sequence conservation in this domain is relatively poor, and whereas it seems likely that such a narrow part of the pore affects permeability and conduction across the entire family, additional experiments are needed to prove this hypothesis (84). Interestingly, the critical polar amino acids of the P2X2R (Thr³³⁶, Thr³³⁹, Ser³⁴⁰) are conserved in all the P2X7Rs used in the present study, and a specific mutation of at least one of these (Ser³⁴² of the rat P2X7aR, equivalent to Thr³³⁹ of the rat P2X2R) results in a loss-of-function (95).

Now, our experiments suggest that a third domain should also be considered. Murine P2X7aRs and P2X7kRs have identical pore forming TM2s but divergent $Pf\%$, and differ only in the primary sequences of their N termini and TM1s. Site-directed mutagenesis of the TM1 of the rat P2X2R affects gating (96, 97) to a greater extent than the $Pf\%$ (62), suggesting that TM1 plays little or no role in modulation of the Ca²⁺ component of the ATP-gated current. We assume that this condition holds true for P2X7Rs too, and we plan to test this hypothesis in future experiments. If true, then the N terminus must be the site responsible for the different $Pf\%$ values of the two splice variants. Mutagenesis of the rat P2X7aR N terminus affects the conformational changes that accompany "pore dilation," a process by which a limiting constriction in the pore widens to the extent that large cations like NMDG⁺ show appreciable permeability (65, 95, 98). We suggest that dilation of the pore results in disruption of the putative Domain 2 Ca²⁺ selectivity filter. In this scenario, Domain 2 is made of the side chains (or backbone carbonyl oxygens) of critical polar amino acids (for example, Ser³³⁹ and Ser³⁴² of the rat P2X7aR) that loosely coordinate partially dehydrated Ca²⁺ and thereby facilitate Ca²⁺ transport through the pore. This binding site is disrupted in the constitutively dilated P2X7kRs where the distance between the polar amino acids is too great to provide the electrostatic potential needed to selectively bind Ca²⁺ at the expense of Na⁺ and K⁺. In 2004, Fisher *et al.* (99) speculated that pore dilation could alter the preferential flow of one ion over another, and our results support this proposition.

This said, we cannot rule out the possibility that pore dilation and regulation of the Ca²⁺ current are unrelated processes controlled by distinct domains, and that the co-occurrence of constitutive pore dilation and limited Ca²⁺ flux in the P2X7kR is coincidental. One way to address this issue is to study both processes in a single cell before and after onset of pore dilation, which might be possible for cells expressing the rat P2X7aR. Unfortunately, significant pore dilation measured in physiological concentrations of divalent cations requires a prolonged agonist exposure that results in a progressive leak of fura-2 out of the cell and Ca²⁺ into the cell (98); the unintended result is a saturation of the remaining fluorescent dye that negates the accuracy of the $Pf\%$ measurement. Future experiments on mutant P2X7aRs with altered pore properties may help to determine whether one or more structural domains underlie genesis of these two effects.

In conclusion, we show that P2X7aRs transduce a significant Ca²⁺ flux that is as large or larger than those of most ligand-gated channel families. In contrast, the constitutively dilated P2X7kRs show no selective preference for Ca²⁺, which most likely reflects an N-terminal directed change in the conformation of the conducting pore. The results bring to light the intriguing hypothesis that the physiological consequence of pore dilation is a decrease in Ca²⁺ entry (99). Such a mechanism may have evolved to prevent Ca²⁺ overload in cells expressing P2X7aRs that show a time-dependent facilitation of peak current amplitude, or in cells expressing P2X7kRs with altered channel kinetics and/or increased sensitivities to ATP. Our data also demonstrate an important and somewhat overlooked physiological consequence of the substantial buffering

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of Ca²⁺ at the high [ATP]_o values that accompany disease. That is, elevated levels of extracellular ATP lead to smaller than expected Ca²⁺ currents, which would blunt agonist-gated responses triggered by an elevation of intracellular [Ca²⁺]. At the same time, the reduction of [Ca²⁺]_o by ATP would partially relieve the allosteric Ca²⁺ block of the P2X7R, leading to a larger than expected peak current. In the future, it will be important to consider all of these factors when attempting to gauge the likely response of any cell expressing a P2X7R to ATP.

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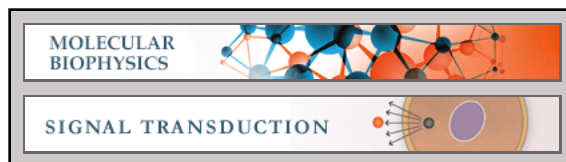
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