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P2X2 and P2X5 Subunits Define a New Heteromeric Receptor with P2X7-Like Properties

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Ligand-gated ion channels are prototypic oligomeric membrane proteins whose stoichiometry determines their functional properties and subcellular localization. Deciphering the quaternary structure of such protein complexes is an arduous task and usually requires the combination of multiple approaches. ATP-gated P2X receptors are formed by the association of three subunits, but the quaternary arrangement of the seven P2X subunits at the plasma membrane remains poorly characterized. By combining bioluminescence resonance energy transfer, bifunctional fluorescence complementation and protein biochemistry, we developed an experimental approach that allows precise determination of rat P2X receptor quaternary assembly. We found that P2X5 subunits associate with P2X1, P2X2, and P2X4 subunits. We demonstrate that P2X5 and P2X2 subunits interact to form as yet uncharacterized heteromeric receptors with alternate stoichiometries, both present at the plasma membrane. P2X2/5 receptors display functional properties such as pore dilatation, membrane blebbing, and phosphatidylserine exposure that were previously thought to be characteristic hallmarks of the P2X7 receptor. In mouse, P2X2 and P2X5 subunits colocalize and physically interact in specific neuronal populations suggesting that other P2X receptors might contribute to cellular responses typically attributed to P2X7 receptor.

Introduction

Ligand-gated channels are prototypic oligomeric membrane proteins formed by the association of a given number of the same subunit (homomeric channels) or of different subunits (heteromeric channels) (Taly et al., 2009). The subunit composition of ligand-gated channels defines not only their pharmacological properties but also their functional characteristics. However, deciphering the quaternary subunit assembly of channels can sometimes lead to controversial results.

P2X receptors are ATP-gated channels with widespread tissue distribution and physiological functions (Surprenant and North, 2009). Seven different P2X subunits (P2X1–7) have been identified; they all share the same membrane topology with two transmembrane segments linked by a large ectodomain, and the N and C termini are intracellular. Early studies, confirmed by the resolution of the crystal structure of a P2X receptor, proposed that three subunits assemble to form P2X receptors (Nicke et al., 1998; Kawate et al., 2009). Of the seven rat recombinant P2X subunits, five assemble to form functional homomeric receptors in heterologous expression systems, whereas P2X5 receptors are poorly

functional and P2X6 subunits are unable to self-associate (North and Surprenant, 2000; Roberts et al., 2006). Different P2X subunits can also associate as heteromeric receptors. Based on coexpression of recombinant subunits, 13 potential heteromeric P2X receptors have been proposed (Torres et al., 1999), some being expressed at the plasma membrane (Aschrafi et al., 2004). However, to date, P2X2/3 and P2X1/5 receptors are the only unambiguously characterized heteromeric P2X receptors in native tissue (Brederson and Jarvis, 2008). The P2X2/3 receptor was initially identified in sensory neurons through its unique pharmacological and biophysical profiles, which were recapitulated in a recombinant system (Lewis et al., 1995; Radford et al., 1997). Similarly, in submucosal arterioles and cortical astrocytes, ATP-induced currents display the same biophysical and pharmacological profiles that are observed in HEK cells cotransfected with P2X1 and P2X5 subunits (Haines et al., 1999; Lê et al., 1999; Surprenant et al., 2000; Lalo et al., 2008).

As with other ligand-gated channels, the experimental strategies required to investigate P2X subunit assembly should lead to the clear determination of the nature of the interactions between subunits and/or receptors, to the exact definition of the stoichiometry of these interactions, and ultimately provide a demonstration of the existence of these interactions in native conditions.

In this study, we used a combination of cell surface assay, bioluminescence resonance energy transfer (BRET), bimolecular fluorescence complementation (BiFC), and biochemical approaches to analyze the interactions existing between P2X subunits. We found that P2X5 subunits can interact with P2X1, P2X2, or P2X4 subunits at the plasma membrane. We demonstrate that P2X2 and P2X5 subunits associate in a new heterotri-

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meric receptor showing two alternate stoichiometries. We provide evidence that P2X2/5 receptors are expressed in specific regions of the CNS, and that P2X2/5 receptors have functional properties that were previously thought to be unique to the P2X7 receptor.

Materials and Methods

cDNA cloning, epitope tagging, and site-directed mutagenesis. The different P2X cDNAs carrying extracellular tags and mutations were described previously (Chaumont et al., 2004). The hemi-yellow fluorescent protein (YFP-N and YFP-C), RAMP1-YC, and CRLR-YN were gifts from Dr. M. Bouvier (University of Montreal, Montreal, Canada) (Héroux et al., 2007); YC3.1 (CAM) was a gift from Dr. B. Khakh (University of California Los Angeles, Los Angeles, CA) (Richler et al., 2008); and *Renilla* luciferase (Rluc) and PAR-1-YFP were gifts from Dr. Ayoub (Institut de Genomique Fonctionnelle, CNRS, Montpellier, France) (Ayoub et al., 2007). All reporter proteins were PCR inserted in frame at the C-terminal tail of P2X subunits.

Cell culture and transfections. Human Embryonic Kidney 293 cells (HEK) were maintained as described previously (Chaumont et al., 2004). Transfections were performed using Lipofectamine 2000 (Invitrogen) according to the manufacturer's recommendation; 4 h after transfection, cells were split and incubated at 37°C for 36–48 h in DMEM supplemented with glutamine, fetal calf serum, and antibiotics.

Bioluminescence resonance energy transfer and bimolecular fluorescence complementation. Transfected HEK cells were plated in two independent 96-well polyornithine-coated plates. Thirty-six hours later, cells were washed twice with calcium and magnesium-supplemented PBS, and coelenterazine H (Invitrogen) was added at a final concentration of 5 μ M. Emitted light was collected at 2 Hz using a Mithras LB 940 plate reader (Berthold Technologies) using two filter settings (Rluc filter, 485 \pm 20 nm; and YFP filter, 530 \pm 25 nm). The BRET signal corresponds to the ratio of YFP/Rluc signals. Values were corrected by subtracting the background BRET signal measured in the absence of YFP constructs. For BRET titration curve experiments, cells were transfected with a fixed amount of Rluc fusion cDNA and increasing amounts of YFP fusion cDNA. To control for relative expression levels of donor and acceptor proteins, total luminescence and fluorescence were measured independently in separate plates. BRET values were then plotted as a function of the total fluorescence/luminescence ratios (YFP/Rluc). For kinetic studies, after baseline acquisition, 100 μ M ATP was applied to individual wells through the Mithras automatic injection system. For BRET/BiFC experiments, cells were incubated at 30°C for 12 h before the reading to promote the complementation between hemi-YFP. Total fluorescence, BRET signal, and YFP recomplementation were measured as described above.

Chemiluminescent assay. ELISA assay was performed as described by Chaumont et al. (2004). Briefly, transfected cells were PFA-fixed and Triton-permeabilized (or not), blocked with PBS with 1% fetal calf serum, and incubated for 30 min with anti-Flag M2 antibody directly coupled to HRP (1/4000) (Sigma). Luminescence was measured using Supersignal femto substrate (Pierce) and quantified in a Victor 2 luminometer (PerkinElmer). Surface expression was calculated as the ratio of the signal obtained in nonpermeabilized cells to the signal obtained for permeabilized cells.

Plasma membrane protein biotinylation, immunoprecipitation, and Western blotting. Transfected HEK cells were washed twice in ice-cold PBS containing 1 mM CaCl₂ and 0.5 mM MgCl₂ (PBS-CM). Cell surface proteins were labeled using 1 mg/ml sulfo-NHS-LC-Biotin (Pierce) for 30 min at 4°C in biotinylation buffer (in mM: 10 H₃BO₃, 140 NaCl, pH 8.8). Cell lysis was performed for 30 min at 4°C under agitation in lysis buffer [in mM: 20 HEPES, 100 NaCl, 5 EDTA, 1% NP40, pH 7.4, and protease inhibitors mixture (Roche Diagnostic)]. Lysates were centrifuged (16,000 \times g, 10 min, 4°C) and the supernatant was collected. For immunoprecipitation, 500 μ g of protein was incubated on a rotating wheel with either anti-HA (12CA5) or anti-Myc (9E10) antibodies and then with protA-Sepharose beads (4°C, 1 h). After washes in lysis buffer, bound proteins were eluted by two incubations with 200 μ g/ml Myc or

HA peptides. Proteins were resolved by SDS-PAGE, transferred to nitrocellulose membranes and visualized using either streptavidin–HRP (1/50,000, Pierce) or appropriate primary antibodies: HA 1/2000 (clone 12CA5, Roche), Myc 1/5000 (clone 9E10, Santa Cruz Biotechnology), GFP 1/2500 (Torrey Pines), EE 1/2000 (Bethyl Laboratories), and Super-Signal West Pico substrate (Pierce).

Cross-linking in vivo of plasma membrane protein. Cross-linking was performed as described above for plasma membrane protein biotinylation, except that BS3 (Pierce) was used at a final concentration of 2 mM. After cross-linking, cells were lysed and proteins separated by SDS/PAGE. Cross-linked P2X subunits were revealed by Western blotting using anti-Myc or anti-GFP antibodies.

PFO-PAGE. Native gels were performed as described previously (Ramjeesingh et al., 1999). Membrane solubilization was performed in lysis buffer as described above for Western blotting, and 2 \times loading buffer was added [100 mM Tris base, 1% (w/v) NaPFO, 20% (v/v) glycerol, 0.005% Bromophenol Blue; pH 8.0 with NaOH]. Electrophoresis was performed at 4°C on polyacrylamide gel in PFO containing running buffer [25 mM Tris, 192 mM glycine and 0.5% (w/v) PFO; pH 8.5 with NaOH]. P2X subunits were revealed as described above. Molecular weight markers apoferritin (440 kDa), β -amylase (200 kDa), alcohol dehydrogenase (150 kDa), and bovine serum albumin (66 kDa) were diluted in 2 \times loading buffer containing NaPFO and separated by electrophoresis as described above. Relative migration of protein standards was detected by Ponceau red staining before membrane blocking.

Coimmunoprecipitation of native P2X5 and P2X2 proteins. Immunoprecipitation of native P2X subunits was performed from membrane protein extracts as previously described (Chaumont et al., 2008). After preclearing, solubilized membrane proteins were incubated overnight at 4°C with either rabbit anti-P2X5 [a gift from Dr. Mark Voigt (Schwiebert et al., 2002)] or rabbit anti-P2X2 (Alomone Labs), and incubated 1 h at 4°C with 20 μ l of proteinA-Sepharose beads. Beads were washed 3 \times 5 min with solubilization buffer, and bound proteins were eluted with Laemmli sample buffer. Immunoprecipitates were detected by Western blotting as described above, except that Rabbit TrueBlot (eBioscience) secondary antibody was used for detection of P2X5, according to the manufacturer's instruction. Anti-P2X2 and anti-P2X5 were used at 1/500 to reveal the corresponding proteins.

Immunocytochemistry on HEK cells. Transfected HEK cells were plated on polyornithine-coated 25 mm glass coverslips and maintained for 2–3 d in culture. For *in vivo* labeling of tagged receptors, cells were incubated directly in the culture medium containing anti-Myc antibody (9E10, 1/1000), for 45 min at 37°C. After washes, cells were fixed in 4% paraformaldehyde, 4% sucrose PBS-CM for 5 min, washed with a solution of 0.1 M glycine in PBS-CM, and incubated with fluorescent secondary antibody for 45 min. After extensive washing, coverslips were mounted and viewed with a Leica DMRA2 fluorescent microscope.

Immunocytochemistry on native tissue. For experiments on dorsal root ganglia (DRGs), mice of either sex were anesthetized with pentobarbital (300 mg/kg), and DRGs from L6, L5, L4 were dissected and fixed 2 h in 4% paraformaldehyde. For spinal cord and brainstem, mice were transcardially perfused with 4% paraformaldehyde; postfixation was performed in the same solution at 4°C overnight. Tissue sections (40 μ m) were made with a vibratome, washed with PBS, and blocked for 1 h in PBS containing 10% BSA, 0.1% Triton X100 with guinea pig anti-P2X2 (1/1000) (NeuroMics), and rabbit anti-P2X5 (1/500). After washing, slices were incubated for 4 h with the corresponding secondary antibody. After washing, sections were mounted in mowiol (Calbiochem). Images were acquired on a Biorad MRC 1024 laser-scanning confocal microscope.

Video-microscopy. Time-lapse video-microscopy of membrane blebbing was performed on transfected HEK cells plated on 35 mm glass-bottom dishes (Fluorodish, WPI) precoated with polyornithine. Experiments were performed in DMEM-HEPES culture medium in a temperature-regulated chamber at 37°C using a Zeiss Axiovert 2000M equipped with a Cool Snap Roper Scientific HQ camera and a \times 63 objective. Image acquisition started 2 min before perfusion with 100 μ M ATP and lasted for 5 min, with an image acquisition rate of 0.5 Hz.

Dye uptake. Yo-Pro-1 uptake: Transfected HEK cells were plated as described above. Cells were washed and then incubated for 5 min before video-microscopy in NMDG medium (in mM: 147 NMDG, 0.3 CaCl₂, 2 KCl, 1 MgCl₂, 10 HEPES, 12 glucose, pH 7.4) containing 2 μM Yo-Pro-1 (Invitrogen). Some experiments were performed in physiological medium containing 147 mM NaCl instead of NMDG. The Metafluor Imaging system (Molecular Devices) was used for fluorescence acquisition and analysis of individual cells. After 30 s of baseline acquisition, cells were perfused with 10 or 100 μM ATP for the following 3 min. Images were acquired every 300 ms except where indicated. Fluorescence was excited at 480 nm by illumination through ×20 water-immersion objective and detected above 510 nm with a CCD camera. Results are expressed as the mean of all recorded cells (>50) after background subtraction. Ethidium bromide (EtBr) uptake experiments were performed as described above except that fluorescence was detected at 590 nm after excitation at 540 nm. Cells incubated in NaCl-based solution and EtBr (25 μM) were stimulated with 100 μM or 5 mM ATP. Images were acquired every 20 s using a Zeiss Axiovert 25 microscope equipped with a Zeiss AxioCam HRc camera. Fluorescence variations were quantified using ImageJ software (<http://rsb.info.nih.gov/ij/>).

Annexin-V staining. P2X-YFP transfected HEK cells seeded on polyornithine-coated glass coverslips were stimulated with the appropriate ATP concentration for 2 min in normal culture medium, rinsed with PBS, and incubated for 5 min with annexin-V-Cy3 according to the manufacturer's instruction (Biovision). After washing, cells were fixed and the coverslips mounted and viewed with a Leica DMRA2 fluorescent microscope.

Electrophysiology. For *Xenopus laevis* oocyte experiments, rat P2X2 and P2X5 cDNAs were linearized and *in vitro* transcribed using SP6 polymerases (Mmessage-machine; Ambion). Final RNA concentration was adjusted to 1 μg/μl. A 1:3 mixture of P2X2 and P2X5 cRNAs was used to favor the expression of P2X5 subunits over that of P2X2. P2X2 cRNA was diluted with water accordingly. cRNA was injected into freshly isolated *Xenopus* oocytes. Two to three days after injection, currents were recorded under two-electrode voltage-clamp using a GeneClamp 500 amplifier (Molecular Devices) and digitized at 500 Hz. Current and voltage electrodes (1–1.5 MΩ) were filled with 3 M KCl, and the membrane potential was clamped at –60 mV. Oocytes were perfused at a flow rate of 5 ml/min with ND96 solution containing in mM: 98 NaCl, 2 KCl, 1.8 CaCl₂, 1 MgCl₂, 5 HEPES, pH 7.4. NMDG recording solution contained in mM: 98 NMDG, 2 KCl, 2 MgCl₂, 5 HEPES, pH 7.4. All drugs (Sigma-Aldrich) were applied at their final concentration using a homemade computer-driven valve system. Current–voltage relations were obtained by applying 200 ms voltage ramps from –100 to +60 mV at 1 Hz; reversal potentials were determined from the *I*–*V* relationship using Clampfit software. Dose–response curves were fitted to the Hill sigmoid equation, and EC₅₀ values were determined by nonlinear regression analysis (Prism, GraphPad). Statistical comparisons were

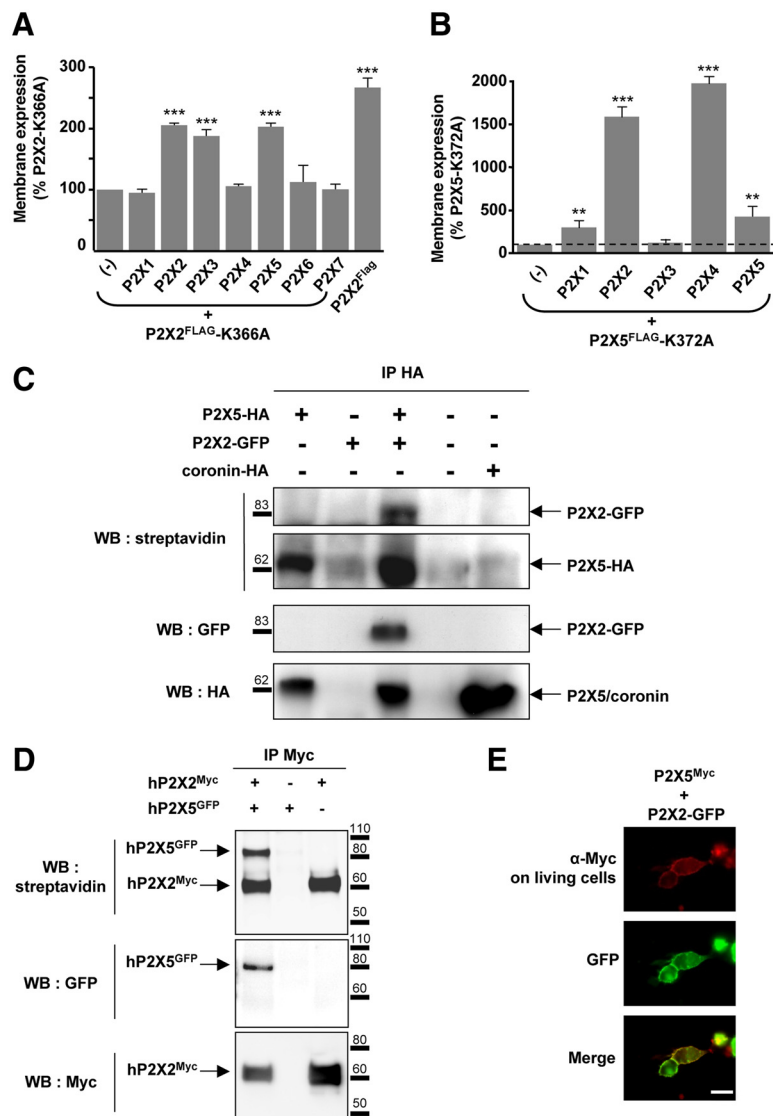


Figure 1. P2X5 subunits interact with P2X2 subunits at the plasma membrane. **A**, Rescue of cell surface expression of the P2X2–K366A trafficking deficient mutant by other subunits. P2X2–K366A carrying a Flag tag in the extracellular loop was expressed alone or in combination with each of the seven wild-type P2X subunits. Membrane expression was measured using a chemiluminescent assay. Cell surface expression of P2X2–K366A is rescued upon coexpression with P2X2, P2X3, P2X4, and P2X5 subunits. **B**, Membrane detection of the P2X5–K372A mutant carrying an extracellular Flag tag is increased when coexpressed with P2X1, P2X2, and P2X4 and P2X5 subunits. Note that because the surface expression of P2X5–K372 mutant is very low compared with P2X2–K366A, the changes appear much higher. Results are shown as mean \pm SEM of at least three independent experiments. $^{**}p < 0.01$, $^{***}p < 0.005$, Student's *t* test. **C**, P2X5 subunits interact with P2X2 at the plasma membrane. P2X5 carrying an extracellular HA tag was expressed alone or in combination with P2X2–GFP. Immunoprecipitation was performed after labeling living cells with sulfo-NHS-LC-biotin. Top, Biotinylated protein fraction after HA immunoprecipitation detected with streptavidin–HRP. Middle and bottom, Detection of coimmunoprecipitated GFP and HA tagged-proteins, respectively. Coronin was used as a control for cell permeabilization. **D**, Human P2X2 and P2X5 subunits interact at the plasma membrane. Experiments were carried as described above, except that human P2X2–Myc and human P2X5–GFP cDNAs were used. **E**, Immunodetection of P2X5 in living cells. HEK cells were transfected with P2X5^{Myc} subunits carrying an extracellular Myc tag and P2X2–GFP. Immunostaining of P2X5^{Myc} subunits (red) was performed by incubation of Myc antibodies on living cells; P2X2 subunits (green) were revealed through GFP fluorescence. Scale bar, 10 μm.

assessed using Student's *t* test. The differences were considered significant for $p < 0.05$.

Whole-cell recordings of HEK cells were performed at room temperature 24–48 h after transfection, with a patch-clamp amplifier (Axopatch 200B) using pCLAMP9 software (Molecular Devices), at a holding potential of –60 mV. All data were analyzed using pCLAMP9 and GraphPad Prism (GraphPad) software.

For rescue of P2X trafficking deficient subunits, the extracellular solution contained in mM: 147 NaCl, 2 KCl, 2 CaCl₂, 1 MgCl₂, 10 HEPES,

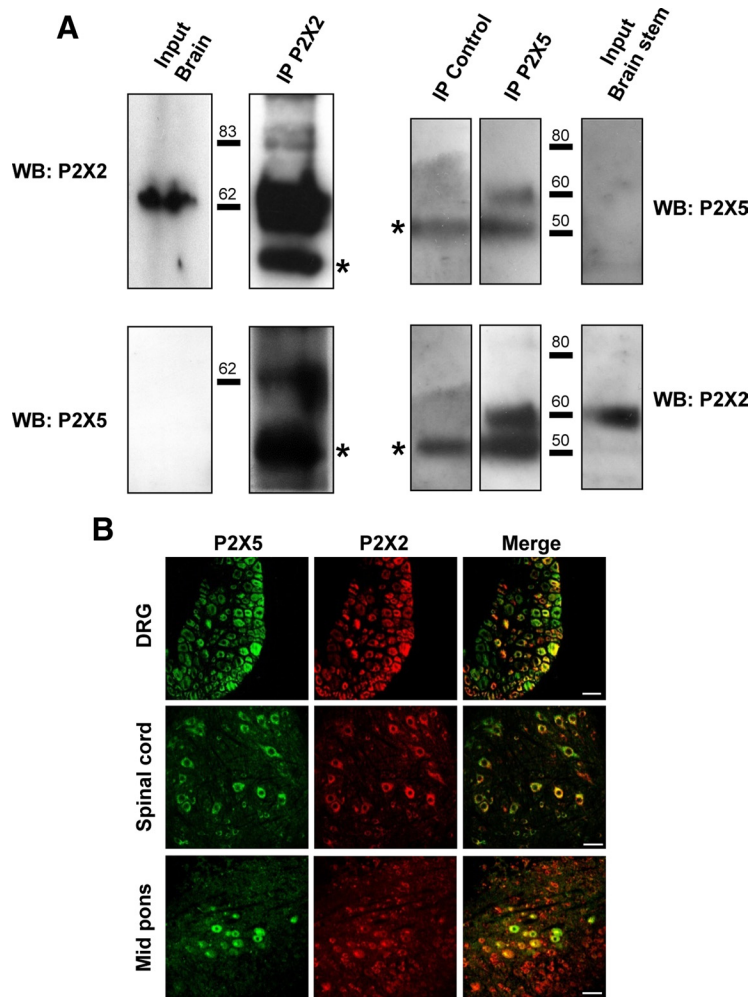


Figure 2. P2X5 and P2X2 subunits interact in neuronal tissues. **A**, Physical interactions of P2X2 and P2X5 subunits in brain tissues. P2X2 or P2X5 subunits were immunoprecipitated from mouse brain membrane and different subunits detected with specific antibodies. Left panel represents immunoprecipitation from total brain membrane proteins, right panel from brainstem. Note that P2X5 is not detectable from the total protein extract. Blots are representative of three independent experiments. Asterisks indicate the antibody heavy chains. **B**, Coimmunolocalization of P2X2 and P2X5 subunits in peripheral and central neurons. Representative images of immunohistochemistry performed on slices from DRG, spinal cord, or mid pons. P2X2 and P2X5 are stained in red and green, respectively. Note that in each structure only a subset of neurons coexpresses both subunits. Scale bar, 40 μm .

and 13 D-glucose, and the intracellular pipette solution contained in mM: 145 NaCl, 10 HEPES, and 10 EGTA (pH adjusted to 7.3 with NaOH, ~ 320 mOsm/L). Data were low-pass filtered at 2 kHz and digitized at 5 kHz. Agonists were applied using an RSC 200 system (Biological Science Instruments). Test concentrations of ATP and BzATP (100 and 300 μM , respectively) were applied for 2 s at 2 min intervals, alternating which agonist was applied first.

For NMDG permeability experiments, the extracellular solution contained in mM either: 135 NaCl, 20 TEACl, 2 CaCl₂, 1 MgCl₂, and 10 HEPES (pH adjusted to 7.3 with KOH, ~ 320 mOsm) or 160 NMDG, 0.1 CaCl₂, 1 MgCl₂, and 10 HEPES (pH adjusted to 7.3 with HCl). Borosilicate glass pipettes had a typical resistance of 1.5–2.5 M Ω when filled with an internal solution containing in mM: 140 CsCl, 10 EGTA, 10 HEPES, 3 Mg-ATP, 0.6 GTPNa, and 3 CaCl₂ (pH adjusted to 7.3 with KOH, ~ 315 mOsm). Junction potentials in NMDG solutions (~ 10 mV) were not compensated. Drugs were applied by a homemade gravity-driven perfusion system.

Results

Determination of P2X subunit interactions at the plasma membrane

Using a cellular ELISA assay, we previously demonstrated that membrane expression of trafficking-deficient mutants of rat P2X

subunits could be rescued by coexpression with wild-type subunits, likely within a heteromeric channel (Chaumont et al., 2004). We used this property in the present study to screen for any rat subunits potentially interacting with P2X2. Our results indicated that in addition to P2X2 and P2X3, only the P2X5 subunit was able to rescue membrane expression of the P2X2-K366A trafficking mutant (Fig. 1A). We next performed the converse experiment to investigate subunits that might interact with rat P2X5. As shown in Figure 1B, membrane expression of the P2X5-K372A trafficking mutant was rescued upon coexpression with P2X1, P2X2, P2X4, and P2X5 subunits. In this case, because the signal of P2X5-K372A at the plasma membrane was very low (data not shown), the level of rescue of P2X5-mutated subunits appeared substantially higher than observed for P2X2-K366A. According to our interpretation that membrane expression of trafficking-deficient subunits can be rescued by specific interaction with any wild-type subunit, these results suggest that P2X5 subunits can specifically assemble with P2X1, P2X2, P2X4, and P2X5, establishing P2X2 and P2X4 as new potential partners of P2X5. P2X4 interactions with P2X5 were not investigated further in this study.

To confirm the interaction between P2X2 and P2X5 subunits, we performed immunoprecipitation after biotinylation of membrane proteins in living cells. HEK cells were transfected with P2X5 carrying a HA tag either alone or in combination with P2X2 subunits carrying a C-terminal GFP. After 48 h, living cells were biotinylated and P2X5 subunits were immunoprecipitated using an anti-HA antibody. The results

shown in Figure 1C indicate that P2X5 coimmunoprecipitates with P2X2 subunits and that all the subunits are biotinylated, establishing that P2X5 and P2X2 subunits interact at the plasma membrane. Identical results were obtained with full-length human P2X2 and P2X5 subunits (Fig. 1D). However, as previously described (Kotnis et al., 2010), we observed that the human P2X5 subunit with exon 10 deleted (hP2X5- ΔX , a form predominantly expressed in humans) was not expressed at the plasma membrane and triggered a strong reduction of cell surface expression of P2X2 upon coexpression (data not shown), suggesting a dominant-negative effect.

The above results were further confirmed by immunocytochemistry. Rat P2X5^{Myc} was transfected into HEK cells either alone or with P2X2-GFP cDNA. Immunodetection of P2X5^{Myc} was performed on living cells by incubation of the anti-Myc antibody in the culture medium, while P2X2 subunits were detected through GFP fluorescence. Figure 1E shows that P2X5^{Myc} immunostaining was clearly localized to the plasma membrane and closely overlapped with the GFP signals. Altogether, our results demonstrate that P2X5 subunits interact at the plasma membrane with P2X2. These results provide evidence for the existence

of interactions between these subunits, but do not provide information as to whether they occur directly within a heteromeric channel, or indirectly between two homomeric receptors.

Association of P2X5 and P2X2 subunits in brain tissues

In situ hybridization data indicates sparse expression of P2X5 subunits within the nervous system, mostly restricted to the DRG, spinal cord, and mid brain nuclei (Collo et al., 1996). We thus performed coimmunoprecipitation of P2X2 and P2X5 from total mouse brain and brainstem membrane protein extracts, respectively. As shown in Figure 2A, both anti-P2X5 and anti-P2X2 antibodies coimmunoprecipitated the alternate subunit. P2X5 protein could not be detected from the total protein extracts, further supporting a low level of expression of this subunit in the CNS. We next analyzed the colocalization of the two subunits by immunohistochemistry. Sections from mouse DRGs, spinal cord, and mid pons were labeled with anti-P2X5 and anti-P2X2 antibodies. In these three regions, clear cellular colocalizations of the two proteins were observed (Fig. 2B). Aside from DRG neurons that showed strong P2X5 immunostaining, P2X5 expression in the spinal cord was restricted to a subpopulation of neurons in lamina VII, and to the trigeminal mesencephalic nucleus in the pons. In all three regions, only a subset of neurons coexpressed both subunits, supporting the specificity of the staining.

P2X subunit interaction allows bioluminescent resonance energy transfer

As stated above, interactions of recombinant P2X5 and P2X2 subunits may occur within a heteromeric channel, or through indirect interactions of homomeric receptors. To answer this question, we used BRET to evaluate whether the distance between P2X5 and P2X2 subunits is compatible with the formation of a heteromeric receptor. P2X2 and P2X5 subunits were fused to either *Renilla* luciferase (Rluc) or enhanced yellow fluorescent protein (YFP) through their C termini. BRET titration curves were performed in living cells for different combinations of subunits: homomeric P2X5 and P2X2 receptors, or in combination of P2X5:P2X2 (Fig. 3). The G-protein-coupled receptor PAR1-YFP and ATP-gated channel P2X4-YFP were used to assess the specificity of BRET signals (Ayoub et al., 2007). As shown in Figure 3A, when P2X5-Rluc was coexpressed with P2X5-YFP, or P2X2-YFP, a specific BRET signal was observed, characterized by a hyperbolic increase of the signal. In converse experiments, specific BRET signals were only observed between P2X2-Rluc and P2X2-YFP or P2X5-YFP but not between P2X2-Rluc and P2X4-YFP or

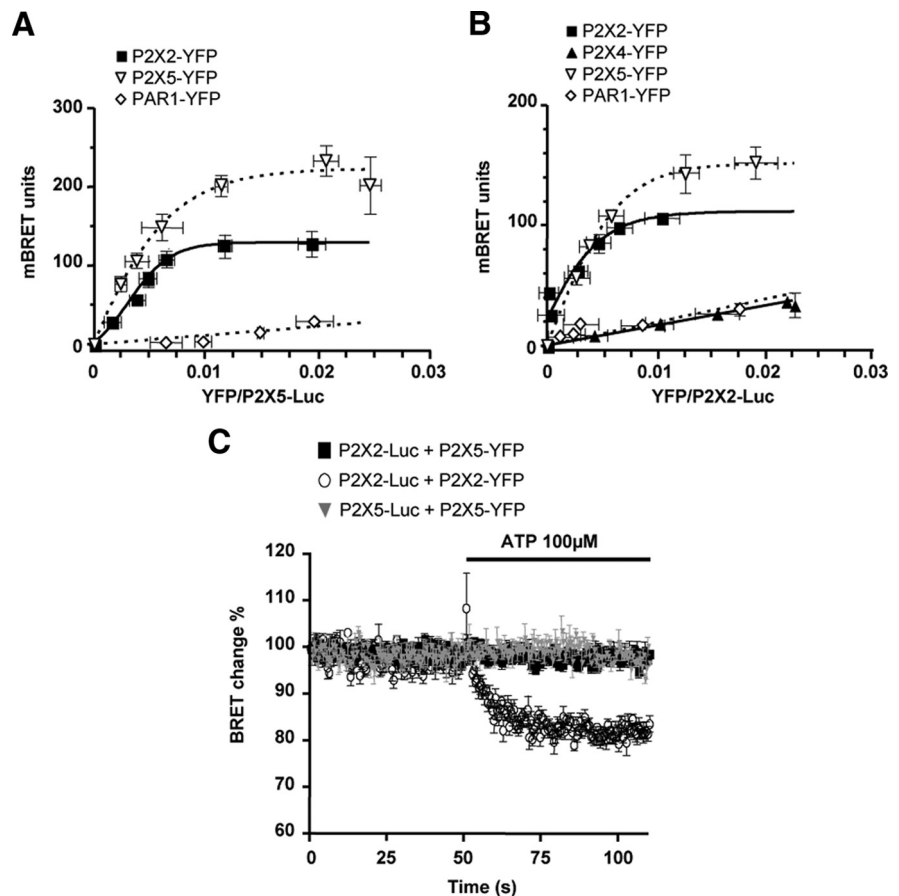


Figure 3. P2X5 subunits are in close spatial proximity to P2X2 subunits. **A**, BRET titration curves using P2X5-Luc as the energy donor. HEK cells were cotransfected with a constant amount of P2X5-Luc and with an increasing amount of YFP fusions. BRET signals are plotted against the relative amounts of each tagged subunit. Specific, saturating, BRET signals are observed between P2X5 and P2X2, or P2X5 subunits, while low linear BRET is obtained with PAR-1. **B**, BRET titration curves using P2X2-Luc as the energy donor, with experiments performed as described above. Specific BRET signals are observed between P2X2 and P2X2 or P2X5 subunits, but not P2X4 and PAR-1. Data are expressed as mean \pm SEM of at least $N = 3$ experiments. **C**, Cytosolic domains of P2X2/5 and P2X2 receptors undergo differential conformational changes after ATP stimulation. Dynamic BRET signals were recorded after ATP stimulation (100 μ M) in HEK cells transfected with P2X2, and P2X5 subunits carrying either a luciferase (Luc) or YFP tag on their C terminus tail, alone or in combination. ATP stimulation induced a diminution of BRET signal between P2X2 subunits, which is not observed for P2X2/5 and P2X5 receptors. Representative experiment reproduced three times. Data are mean \pm SEM of triplicate.

PAR1-YFP (Fig. 3B). These results support the existence of heteromeric P2X2/5 receptors.

The carboxyl cytosolic regions of P2X2 undergo conformational rearrangement that can be evidenced using dynamic measurement of fluorescence energy transfer (FRET) between individual subunits (Fisher et al., 2004). We investigated whether such conformational movements also occurred in P2X2/5 receptors. The dynamics of BRET signals after stimulation with 100 μ M ATP were measured in living HEK cells expressing different combinations of P2X subunits. As shown in Figure 3C, in cells expressing homomeric P2X2 subunits (P2X2-Rluc and P2X2-YFP), stimulation with ATP induced a 20% reduction of the BRET signal, with an estimated time constant of 5 s. In cells expressing homomeric P2X5 receptors or P2X2/5, ATP stimulation did not induce any change in the BRET signal. These results show that ATP-induced conformational movements of the cytosolic tails of P2X2 subunits are either impaired or undetectable when coexpressed with P2X5. Alternatively, these conformational changes might be specific to homomeric P2X2 receptors.

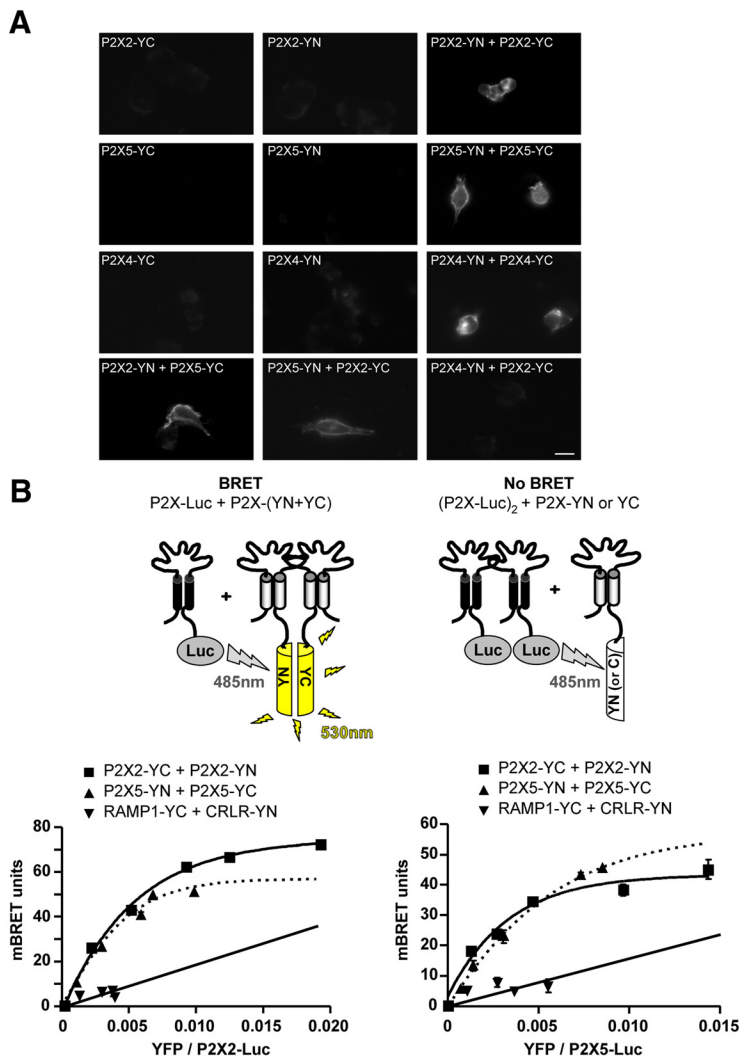


Figure 4. Stoichiometry of P2X2 and P2X5 interactions as assessed by BiFC and BRET/BiFC. **A**, Bimolecular fluorescence complementation between P2X subunits. P2X subunits fused to either the amino or carboxyl half of YFP (YN or YC) were transfected alone or in combination in HEK cells. Recombplemented fluorescence was observed by microscopy. BiFC was observed between homomeric P2X2, P2X4, and P2X5 subunits, as well as between heteromeric P2X2 and P2X5 subunits, but not between P2X2 and P2X4 subunits. Scale bar, 10 μ m. **B**, Combination of BRET and BiFC reveals the stoichiometry of the heteromeric P2X2/P2X5 assembly. Top, Diagram illustrating the approach. BRET is only observed between one P2X2-Luc subunit and any other combinations of two P2X-hemi-YFP subunits. BRET titration curves between P2X2-Luc (left) or P2X5-Luc (right) cotransfected with P2X2, P2X5, RAMP1, and CRL fused to hemi-YFP. Specific BRET signals were observed between P2X2 and P2X5 subunits for each experimental condition, demonstrating the existence of heteromeric receptors with two different stoichiometries. Data are expressed as mean \pm SEM of at least $N = 3$ experiments.

Analysis of P2X2/5 receptor stoichiometry by combined BRET and bimolecular fluorescence complementation

We next sought to determine the stoichiometry of the P2X2/5 complex in living cells by combining BiFC and BRET approaches. We first analyzed BiFC in homomeric P2X subunit assemblies. P2X2, P2X4, and P2X5 fused through their C termini to either the amino or carboxyl half of YFP (YN and YC, respectively) were transfected in HEK cells, either alone or in combination. As shown in Figure 4A, complementation of the YFP fluorescence was observed for the three homomeric combinations of subunits tested. Similarly, BiFC was also observed between P2X2 and P2X5 subunits, but not P2X4 and P2X2 (Fig. 4A). We next probed the subunit stoichiometry of P2X2/5 receptors by combining BiFC and BRET techniques. Cells were cotransfected with [P2X2-Rluc + P2X2-YN + P2X2-YC], [P2X2-Rluc + P2X5-YN + P2X5-YC], [P2X5-Rluc + P2X5-YN + P2X5-YC], or [P2X5-Rluc + P2X2-

YN + P2X2-YC], and BRET titration curves generated. As shown in Figure 4, B and C, specific BRET signals were observed for every combination of subunit tested, thus establishing that heteromeric P2X2/5 display two different stoichiometries in living cells.

Biochemical characterization of P2X2 and P2X5 interactions

The above results support that P2X2 and P2X5 subunits associate in a heteromeric receptor complex. However, energy transfer can result from higher order oligomerization states of membrane receptors (Khakh et al., 2005; Guo et al., 2008). Furthermore, energy transfer experiments do not give an indication as to whether these interactions occur at the plasma membrane or in intracellular compartments. To resolve these issues, biochemical characterization of the P2X2/P2X5 subunit complex was carried out. First, we analyzed the native oligomeric organization of P2X2 and P2X5 subunits using the PFO/PAGE method (Ramjeesingh et al., 1999). We established that 0.5% PFO efficiently solubilized P2X2 and P2X5 receptor complexes into two main forms corresponding to a dimeric and trimeric assembly of subunits, and a minor form corresponding to the monomeric subunit (data not shown).

We next analyzed the quaternary structure of the P2X2/P2X5 complex. Because the molecular weights of P2X2 and P2X5 subunits are similar, we used P2X subunits fused to the YC3.1 camaleon protein (Cam) to obtain good molecular weight discrimination between homomeric and heteromeric receptors. PFO/PAGE analysis shows that the presence of Cam as a fusion partner did not alter the trimeric architecture of the homomeric P2X2 or P2X5 channel complex (Fig. 5A). We next analyzed the homomeric receptor complex made by the association of Cam- and Myc-tagged

subunits (P2X2-Cam:P2X2-Myc and P2X5-Cam:P2X5-Myc). After solubilization and PFO/PAGE, proteins were revealed using an anti-Myc antibody. For both receptors, in addition to the trimeric association of Myc subunits, a single higher molecular weight form was clearly detected (Fig. 5A, B, indicated by an asterisk), which can only correspond to a trimeric receptor made by the association of two Myc-tagged subunits and one Cam tagged. The presence of a single Cam subunit in the trimeric complex is likely due to slower maturation of Cam tagged proteins. Finally, P2X2 and P2X5 heteromeric subunit oligomerization was analyzed. In both conditions (P2X5-Myc:P2X2-Cam or P2X2-Myc:P2X5-Cam), a single high molecular weight form, larger than the homotrimeric Myc subunit oligomer, was detected. These results demonstrate that (1) P2X2 and P2X5 subunits associate within a heterotrimeric receptor complex, and (2) this receptor exists with two different subunit stoichiometries.

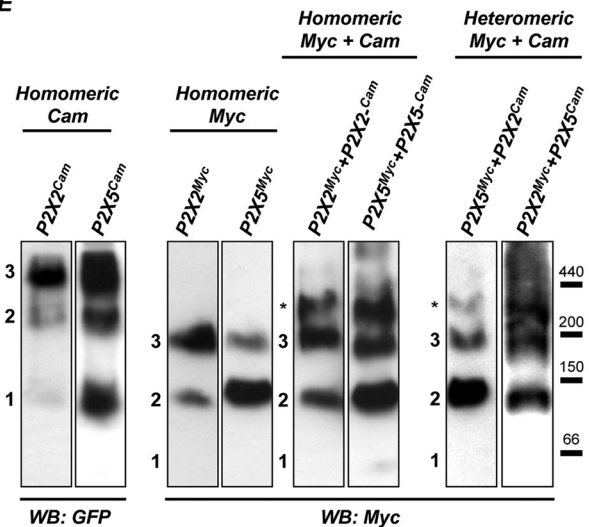
We next asked whether both forms of the heteromeric P2X2/5 receptor are present at the plasma membrane. Plasma membrane P2X receptors were cross-linked in living cells using BS3, a membrane impermeable cross-linker with an 11.4 Å spacer arm, and separated by SDS/PAGE. As shown in Figure 5B, both homotrimeric P2X-Cam and P2X-Myc receptors were efficiently cross-linked at the plasma membrane. However, in the case of P2X-Cam receptors, monomeric subunits were still detected, further supporting that P2X-Cam subunits do not reach the plasma membrane as efficiently as P2X-Myc. Analysis of cross-linked P2X-Cam/P2X-Myc receptors revealed the presence of a single high molecular weight band above the trimeric form of P2X-Myc receptors, which corresponds to a trimeric receptor made by the association of two P2X-Myc subunits and one P2X-Cam subunit. This was true for all combinations of subunits tested (P2X2-Myc + P2X2-Cam, P2X5-Myc + P2X5-Cam, P2X2-Myc + P2X5-Cam, P2X5-Myc + P2X2-Cam). These results establish that the two stoichiometries of P2X2/5 receptors are both expressed at the plasma membrane.

P2X2/5 receptors display specific functional properties

The functional properties of P2X2/5 receptors were analyzed in mRNA-injected *Xenopus* oocytes. To minimize the presence of homomeric P2X2 receptors, P2X2 and P2X5 mRNA were injected at a 1:3 ratio, respectively. The P2X2/5 receptor pharmacological profile was established and compared with that of P2X2 receptors. The results are summarized in Table 1, and show that the apparent affinities of the P2X2/5 receptor for ATP, BzATP, and ATP γ S are somewhat lower than for P2X2 receptors. While ATP and ATP γ S were full agonists for the two receptors, BzATP efficacy was significantly lower at P2X2/5 receptors. Furthermore, TNP-ATP inhibited P2X2/5 to a greater extent than P2X2 receptors, while $\alpha\beta$ meATP was unable to activate either receptor, even at 300 μ M.

Coexpression of P2X2 and P2X5 subunits can lead to a mixture of four different homomeric and heteromeric receptors, making it difficult to precisely characterize the pharmacological profile of heteromeric P2X2/5 receptors. We previously described that, within a heteromeric receptor, function of the P2X2-K366A mutant can be rescued by coexpression with wild-type P2X3 subunits (Chaumont et al., 2004). We sought to determine whether a similar rescue could be obtained upon coexpression with P2X5 subunits. As shown in Table 2, coexpression of P2X5 with P2X2-K366A in

A PFO-PAGE



B BS3 cross-linking

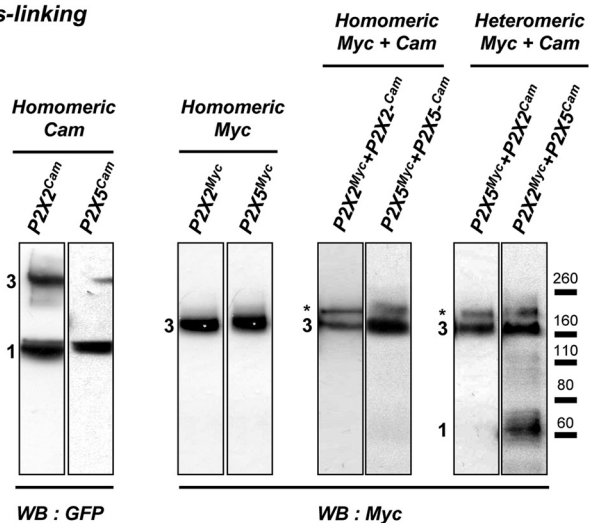


Figure 5. Biochemical analysis of P2X2/5 subunit assembly. **A**, Analysis of the P2X2/5 subunit complex in native conditions using PFO-PAGE. HEK cells were transfected with P2X subunits with either a Myc tag or the Cam protein fused to their C terminus. Homomeric P2X2 and P2X5 receptors fused to Myc or Cam are resolved as monomers, dimers, and trimers (indicated by numbers). When Myc and Cam fused P2X subunits are coexpressed, an additional high molecular weight form of the complex (indicated by an asterisk) is detected, which corresponds to a stoichiometry of two Myc subunits for one Cam subunit. Molecular weight markers apoferritin (440 kDa), β -amylase (200 kDa), alcohol dehydrogenase (150 kDa), and bovine serum albumin (66 kDa) were detected by Ponceau red staining before membrane blocking. **B**, P2X2/5 receptors are present at the plasma membrane with two different stoichiometries. Stoichiometry of P2X receptors at the plasma membrane was analyzed after living cell surface protein cross-linking with the membrane impermeable cross-linker BS3. Cross-linked P2X subunits were analyzed by SDS/PAGE. Homomeric Myc-tagged receptors are resolved as a single band corresponding to the molecular weight of a trimer, while Cam-tagged subunits migrate as monomers and trimers. When Myc- and Cam-fused P2X subunits are coexpressed, an additional high molecular weight form of the cross-linked receptor (indicated by an asterisk) is detected, corresponding to a stoichiometry of one P2X-Cam subunit and two P2X-Myc subunits. All data are representative of $N > 3$ independent experiments.

HEK cells at a ratio of 1:1 yielded currents that were not significantly larger than when either construct were expressed alone. Shifting the ratio of plasmids toward either P2X2-K366A or P2X5 resulted in smaller currents (data not shown). Conversely, coexpression of P2X5-K372A with P2X2 resulted in ATP- and BzATP-evoked currents, with densities similar to cells coexpressing P2X2 and P2X5 subunits. These results suggest that heteromeric (P2X2)₂/P2X5-K372A are functional and display a lower sensitivity to BzATP. However, under these experimental conditions, the presence of functional ho-

Table 1. Pharmacological properties of P2X2, P2X2/5, and P2X5 receptors expressed in *Xenopus* oocytes

	ATP		BzATP		ATP- γ S		TNP-ATP	
	EC50, μ M	I/I_{P2X2} (100 μ M) %	EC50, μ M	I_{BzATP}/I_{ATP} , %	EC50, μ M	$I_{ATP-\gamma S}/I_{ATP}$, %	1 μ M, %	10 μ M, %
P2X2	5.3 \pm 0.9 (<i>n</i> = 5)	100 \pm 0.06 (<i>n</i> = 5)	28 \pm 0.4 (<i>n</i> = 5)	60.6 \pm 6.9 (<i>n</i> = 5)	5.9 \pm 0.2 (<i>n</i> = 6)	94.4 \pm 4.1 (<i>n</i> = 6)	96 \pm 0.9 (<i>n</i> = 3)	46.1 \pm 6.4 (<i>n</i> = 11)
P2X2/5	12.6 \pm 0.4** (<i>n</i> = 13)	90.9 \pm 3.9 (<i>n</i> = 13)	82 \pm 0.9*** (<i>n</i> = 6)	14.1 \pm 3.1*** (<i>n</i> = 6)	22.3 \pm 0.1*** (<i>n</i> = 8)	113 \pm 4.5 (<i>n</i> = 8)	85.9 \pm 3.14 (<i>n</i> = 11)	20.8 \pm 2.5** (<i>n</i> = 11)
P2X5	18.7 \pm 2** (<i>n</i> = 7)	6.7 \pm 2.7*** (<i>n</i> = 7)	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.

In the case of P2X2/5 expression, P2X5 and P2X2 mRNAs were mixed at a 3:1 ratio to minimize the presence of homomeric P2X2 receptors. Consequently, for homomeric P2X2 expression, mRNA was diluted at the same ratio with water. P2X5 mRNA was not diluted for homomeric P2X5 expression. ***p* < 0.01, ****p* < 0.005, unpaired Student's *t* test.

^a300 μ M BzATP, 100 μ M ATP.

^b100 μ M ATP- γ S, 30 μ M ATP.

^cP2X2 was activated with 10 μ M ATP; P2X2/5 was activated with 30 μ M ATP. Percentage of residual current in the presence of TNP-ATP.

Table 2. Analysis of P2X2/5 receptors containing trafficking mutant subunits HEK cells transfected with P2X2, P2X2-K366A, P2X5, and P2X5-K372A subunits or the different combination

	I_{ATP} , pA/pF	I/I_{P2X2} , %	I_{BzATP} , pA/pF	I_{BzATP}/I_{ATP}	<i>n</i>
P2X2	-845 \pm 76	100	-416 \pm 76	0.48 \pm 0.04	18
P2X5	-2.1 \pm 0.9	0.27	—	—	16
P2X2-K366A	-1.5 \pm 0.5	0.15	—	—	6
P2X2-K366A + P2X5	-10.2 \pm 1.6	1.1	-2.8 \pm 0.4	0.35 \pm 0.04*	23
P2X2 + P2X5	-390 \pm 75	45.5	-112 \pm 29	0.26 \pm 0.02**	14
P2X2 + P2X5-K372A	-446 \pm 123	51.6	-104 \pm 25	0.30 \pm 0.03*	9

Currents were evoked by BzATP (300 μ M) and ATP (100 μ M). No response to BzATP was detected for P2X5 or P2X2-K366A. I_{BzATP}/I_{ATP} was calculated using the averaged responses to each agonist from individual cells. Results are shown as mean \pm SEM. **p* < 0.05, ***p* < 0.01 compared to P2X2, ANOVA followed by Tukey–Kramer multiple comparison test.

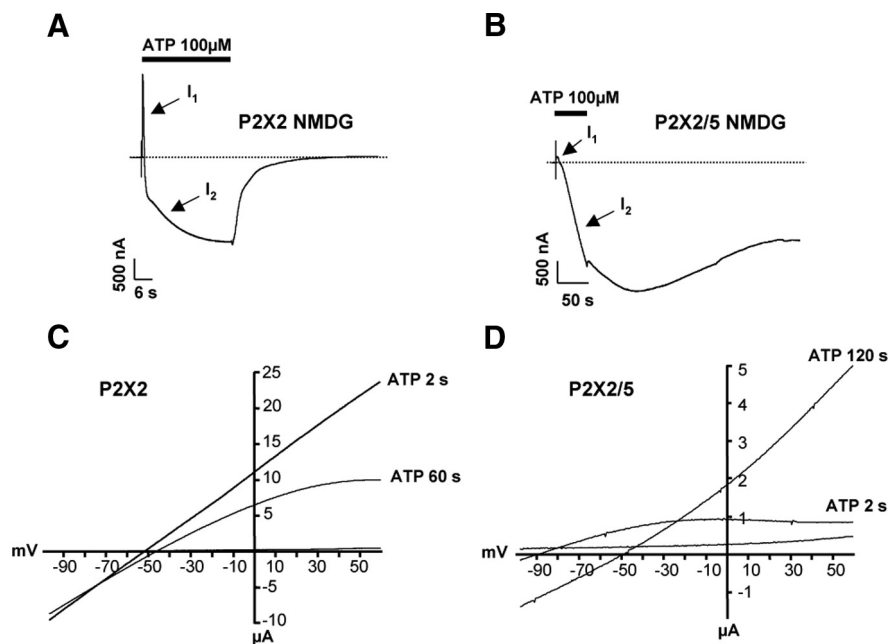


Figure 6. P2X2/5 receptor stimulation evokes sustained NMDG conductance in *Xenopus* oocytes. Permeability dynamics of P2X2 (**A**) and P2X2/5 (**B**). Representative currents evoked by ATP (100 μ M) in *Xenopus* oocytes expressing P2X2 or P2X2/5 receptors recorded in NMDG external solution at a holding potential of -60 mV. I_1 and I_2 correspond to channels permeant to small cations or NMDG⁺, respectively. Note the faster I_1 to I_2 transition for P2X2 receptors compared with P2X2/5, and the lack of reversibility of the inward P2X2/5 conductance. **C, D**, Shift in the reversal potential of ATP-evoked currents in NMDG extracellular solution. Currents were measured during repeated voltage ramps protocol in the presence of 100 μ M ATP. Two seconds after ATP application, reversal potential of P2X2 current has already changed compared with that of P2X2/5. Note the difference in current rectification between the two types of channels.

homomeric P2X2 receptors prevents any precise pharmacological characterization.

P2X2/5 shows functional properties similar to P2X7 receptors

Upon prolonged ATP stimulation, a subset of P2X receptors, including P2X2, can undergo a progressive change in permeabil-

ity, and becoming permeant to large cations such as NMDG (Khakh et al., 1999; Virginio et al., 1999b). Until now, this property has never been investigated in heteromeric receptors. We thus compared the permeability dynamics of P2X2 and P2X2/5 receptors. In P2X2-expressing oocytes, stimulation with 100 μ M ATP in NMDG extracellular solution induced a rapid change in permeability characterized by a transient outward current (I_1) followed by an inward current (I_2), which rapidly reversed upon removal of ATP (Fig. 6A). This change in permeability was also observed as a rapid shift in the reversal potential of the currents. At $t = 2$ s, $E_{rev1} = -72.4 \pm 1.3$ mV (24 cells, 5 experiments), at $t = 60$ s, $E_{rev2} = -57.9 \pm 2.7$ mV (22 cells, 5 experiments). In addition, strong inward rectification gradually developed during

a 60 s ATP application (Fig. 6C). In oocytes expressing P2X2/5 receptors, ATP-induced currents displayed a small initial outward current followed by a slowly developing large inward current (Fig. 6B). The shift in reversal potential was much slower and more pronounced than that observed for P2X2, at $t = 2$ s, $E_{rev1} = -76.4 \pm 2.2$ mV (14 cells, 5 experiments), and at $t = 120$ s, $E_{rev2} = -39.9 \pm 2.3$ mV (14 cells, 5 experiments). The difference of E_{rev2} for P2X2 and P2X2/5-expressing oocytes is significantly different ($p = 0.0058$, Student's *t* test). In contrast to what was observed for P2X2, P2X2/5 currents showed initial strong inward rectification, which progressively disappeared (Fig. 6D). The NMDG conductance persisted after washing off the agonist, eventually returning to baseline; often, the effect was not reversible and led to cell death. This late NMDG conductance was never observed in P2X2-expressing oocytes. Altogether, these results show that the change in permeability observed for P2X2/5 is similar to that of P2X7 receptors.

To further validate these observations, similar experiments were performed in HEK cells in which P2X2/5-evoked conductances were compared with those of P2X7. In HEK cells expressing P2X2/5 receptors, ATP induced a progressive shift in the reversal potential of the currents ($\Delta E_{rev} = 31.3 \pm 3.2$ mV, $n = 9$ cells, two experiments at 60 s), showing a time-dependent increase in NMDG permeability (Fig. 7A, left panel). This shift was more pronounced than in cells expressing P2X7 receptors ($\Delta E_{rev} = 16.8 \pm 1.6$ mV, $n = 6$ cells, 2 experiments) or P2X2

alone ($\Delta E_{rev} = 21.7 \pm 3$ mV, $n = 9$ cells, 2 experiments) (Fig. 7B,C, respectively, left). ΔE_{rev} for P2X2/5- and P2X2-expressing cells were significantly different. As observed in oocytes, a reduction of the outward rectification of P2X2/5 current was observed 60 s after activation, leading to a quasi linear current/voltage curve, as for P2X7-evoked conductances.

We next compared P2X2/5- and P2X7-evoked conductances in NMDG solution. As shown in the right-hand panels of Figure 7, A and B, activation of both receptors evoked currents that persisted after agonist washout. Furthermore, as previously reported (Jiang et al., 2005), this conductance quickly closed after change to an extracellular solution containing NaCl. In contrast, the P2X2-evoked NMDG conductance rapidly closed upon agonist washout (Fig. 7C, right). These data further support that P2X2/5 and P2X7 receptors, but not P2X2, share similar functional properties.

Pore dilatation of P2X receptors allows uptake of fluorescent dyes such as Yo-Pro-1 (Virginio et al., 1999b). We compared the ability of P2X2 and P2X2/5 receptors to mediate Yo-Pro-1 uptake in HEK cells. In NMDG extracellular solution, 100 μ M ATP induced a time-dependent Yo-Pro-1 uptake in cells expressing P2X2 or P2X2/5 (Fig. 8A,B). Compared with P2X2-expressing cells, ATP-evoked Yo-Pro-1 uptake was greater in P2X2/5 transfected cells (Fig. 8C) but had a significantly slower rate [for 100 μ M ATP, $\tau = 1.981 \pm 0.153$ s $^{-1}$ ($N = 4$ experiments) and $\tau = 3.052 \pm 0.133$ s $^{-1}$ ($N = 4$ experiments), for P2X2 and P2X2/5, respectively, $p = 0.039$, Student's t test]. In addition, there was a clear difference in the concentration-dependency of ATP-evoked Yo-Pro-1 uptake between P2X2 and P2X2/5-expressing cells. ATP (10 μ M) induced fluorescence changes of 60 and 25% in cells expressing P2X2 and P2X2/5, respectively (Fig. 8B), while a subsequent stimulation with 100 μ M ATP induced an additional 30 and 75% increase, respectively. Although P2X2/5-mediated Yo-Pro-1 uptake was slower, it evoked a significantly higher final level of fluorescence than was seen with P2X2 receptors (Fig. 8C). Interestingly, for both receptors, Yo-Pro-1 uptake was also observed in physiological solutions, albeit with lower efficacy (Fig. 8D). Similarly, in NaCl-based solution, ATP stimulation evoked EtBr uptake in HEK cells expressing P2X2, P2X2/5, or P2X7 receptors (Fig. 8E). Compared with P2X2, the kinetics of dye uptake was slower and almost linear for P2X2/5 and P2X7. After 2 min, the increase of fluorescence induced by P2X2 reached a plateau, whereas accumulation of dye still occurred 10 min after stimulation in cells expressing P2X2/5 or P2X7. Altogether, these results show that dye uptake induced by P2X2/5 receptors and P2X7 display similar features.

One of the characteristics of the P2X7 receptor is its ability to induce membrane blebbing upon short ATP applications (Mack-

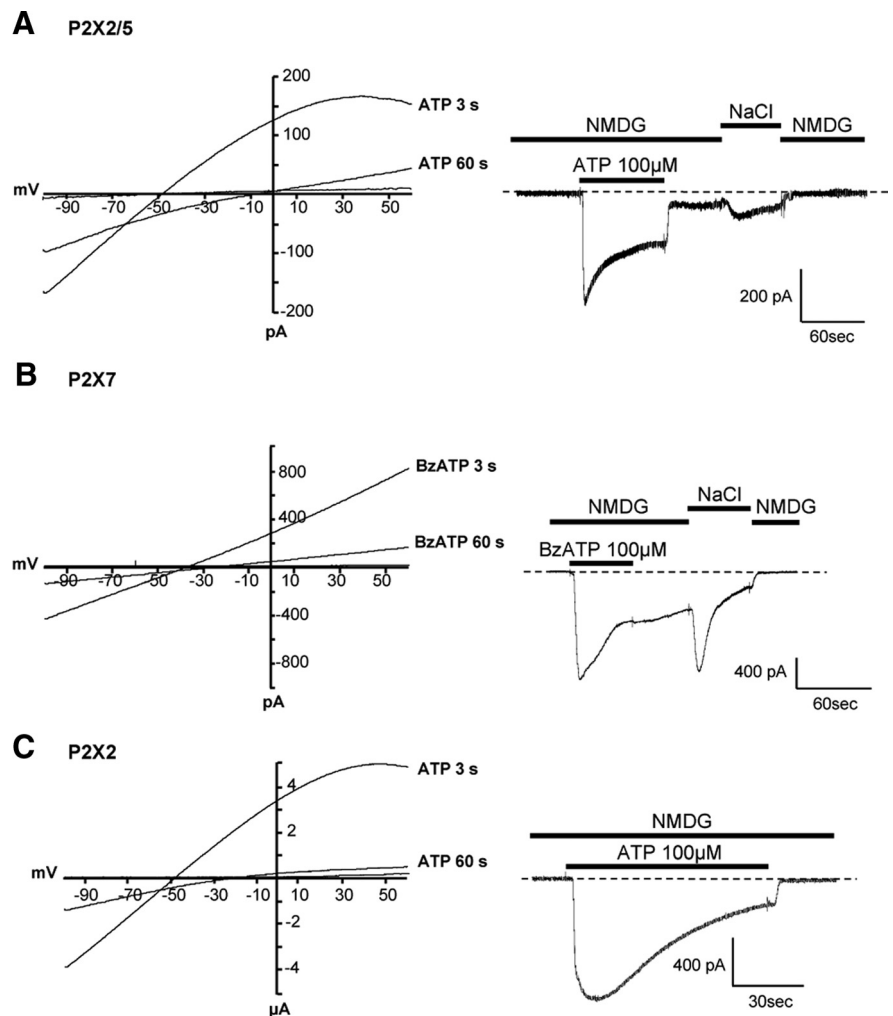


Figure 7. P2X2/5 and P2X7 NMDG conductances display similar properties in HEK cells. Left, Shift in the reversal potential of ATP-evoked currents in NMDG extracellular solution in HEK cells expressing P2X2/5 (A), P2X7 (B), or P2X2 (C). Currents elicited by 100 μ M ATP were recorded during 100 ms voltage ramps from -100 to $+80$ mV, applied from a holding potential of -60 mV. Right, Representative currents recorded in NMDG external solution evoked by 100 μ M ATP (or 100 μ M BzATP for P2X7) in HEK cells expressing P2X2/5 (A), P2X7 (B) or P2X2 (C). Holding potential -60 mV. P2X2/5 and P2X7-expressing cells show a NMDG conductance that persists after agonist washout and can be fully blocked by extracellular NaCl. Note that in P2X2-expressing cells, the NMDG conductance readily reverses upon agonist washout.

enzie et al., 2005; Morelli et al., 2003; Verhoeve et al., 2003). We thus investigated whether P2X2/5 activation could generate a similar phenotype. The membrane dynamics of HEK cells expressing P2X2/5-GFP were recorded using fluorescent time-lapse video-microscopy under normal culture conditions. Application of 100 μ M ATP triggered the appearance of membrane blebs in 50% of transfected cells (Fig. 9A,B). Interestingly, the overall membrane integrity was maintained throughout the 2 min recording period. In similar conditions, there was no difference in the number of blebbing cells between those expressing P2X2-GFP or P2X5-GFP, whereas P2X7-GFP activation induced membrane blebbing in 80% of transfected cells (Fig. 9B).

We next analyzed whether, as for P2X7 receptors, this membrane blebbing was associated with phosphatidylserine exposure (PS flip) and pseudoapoptosis (Mackenzie et al., 2005). HEK cells in physiological culture medium were stimulated with 100 μ M ATP for 5 min, and PS flip was analyzed using annexin-V staining at different times following the stimulation (0, 3, and 24 h). Cells transfected with P2X2-YFP and P2X7-YFP were used as negative

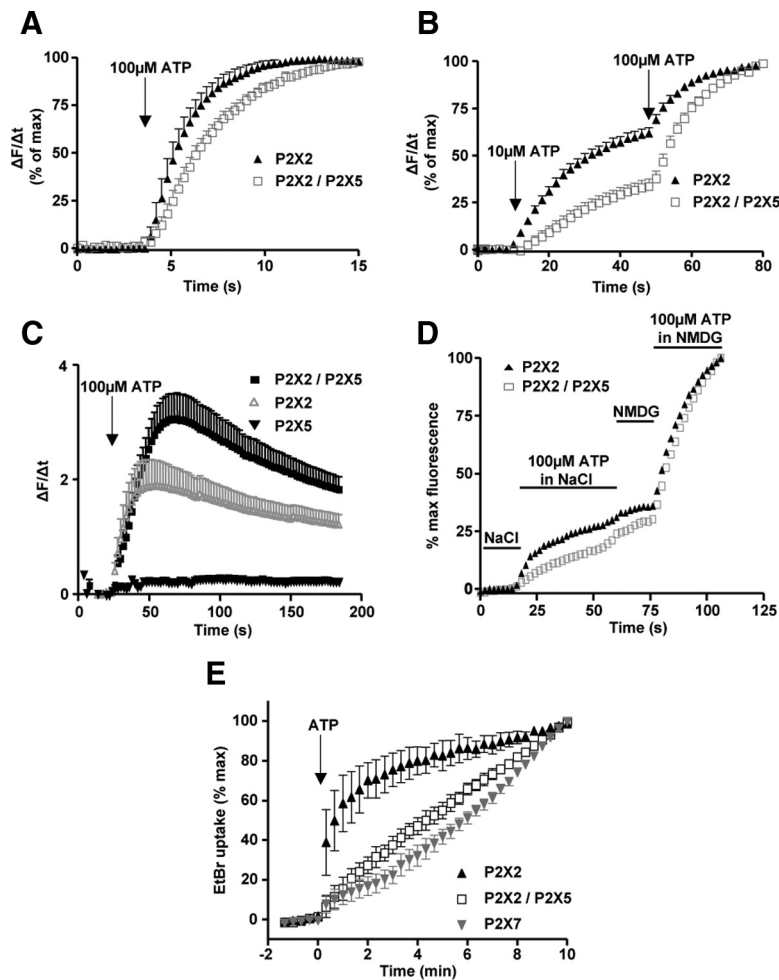


Figure 8. Activation of P2X2/5 receptors induces dye uptake. *A, B*, ATP-evoked Yo-Pro-1 uptake in HEK cells expressing P2X2 or P2X2/5 receptors. Video-microscopy experiments were performed in NMDG extracellular solution. Individual cell fluorescence signals were differentiated and normalized to the maximal response. Data are mean \pm SEM of $n > 10$ cells, $N = 3-4$ experiments. Note that the kinetics of Yo-Pro-1 uptake are slower for P2X2/5 receptor compared with P2X2. *C*, P2X2/5 displays greater ATP-evoked Yo-Pro-1 uptake compared with P2X2. Experiments were performed as above, except that results were not normalized. Data are mean \pm SEM of $n > 10$ cells, $N = 4$ experiments. *D*, ATP-evoked Yo-Pro-1 uptake occurs in physiological solution. Cells were stimulated as above, first in physiological solution (NaCl-based solution) and then in NMDG-based solution. In both conditions, ATP evoked Yo-Pro-1 uptake albeit with higher potency in NMDG solution. Data are mean of $n > 10$ cells, $N = 3$ experiments. SEM was omitted for clarity. *E*, Comparison of ethidium bromide uptake evoked by P2X2, P2X2/5, or P2X7 receptors stimulation. Cells were stimulated with 100 μ M ATP (5 mM for P2X7) in NaCl-based solution. Results are mean \pm SEM, $n > 30$ cells, $N = 4$ experiments. Note that P2X2/5- and P2X7-evoked EtBr uptakes do not saturate.

and positive controls, respectively. In cells expressing P2X2/5-YFP, $>80\%$ of YFP-positive cells also showed annexin-V staining at time 0, and this proportion decreased to $<50\%$ 3 h after the stimulation (Fig. 9*C,D*). After 24 h, the number of annexin-V positive cells was not different from control. A similar trend was observed in cells expressing P2X7-YFP, while in P2X2-YFP transfected cells, annexin-V staining was not different from background. In all conditions, the number of YFP-positive cells remained unchanged, indicating that PS flip induced by short ATP applications did not engage cells in apoptotic cell death.

Discussion

Two main results have emerged from this work. First, we have established that the quaternary organization of P2X subunits can be accurately defined using a combination of BRET and fluorescence complementation. Second, we have demonstrated that P2X5 and P2X2 subunits associate in a new heteromeric receptor with two

stoichiometries. We have provided evidence that this receptor is endogenously expressed within specific neuronal populations, and displays functional properties that were previously thought to be unique to the P2X7 receptor (Surprenant et al., 1996).

Our cell surface assay (Chaumont et al., 2004) allowed a preliminary screening of P2X subunits interacting at the plasma membrane. Indeed, this approach has accurately identified subunits known to interact with P2X2 or P2X5, such as P2X3 and P2X1, respectively (Lewis et al., 1995; Haines et al., 1999; Lè et al., 1999; Surprenant et al., 2000), and revealed P2X2 subunits as a new assembly partner of P2X5. In native tissue, P2X2 and P2X5 subunits were coimmunoprecipitated from brain and brainstem, and immunostaining colocalized the two proteins within specific subsets of neurons known to express P2X5 subunits (Collo et al., 1996). Interactions between P2X5 and P2X2 subunits enable bioluminescent resonance energy transfer, as well as hemi-YFP complementation and BRET/BiFC. These results strongly support that P2X2 subunits associate with P2X5 within a heterotrimeric channel. It is unlikely that the energy transfer observed between these subunits results from nonspecific interactions since BRET signals between P2X2 and P2X5 were clearly saturable, and in parallel experiments there was no energy transfer between P2X2 and P2X4 subunits. Nevertheless, both energy transfer and fluorescence complementation could result from higher state oligomerization of receptor complexes, as demonstrated in the case of dopamine D₂ receptors (Guo et al., 2008). Such FRET-compatible oligomeric interactions have also been described between P2X receptors and different ligand-gated channels, for example between P2X2 and nicotinic channels (Khakh et al., 2005) or between P2X5 and ASIC3 (Birdsong et al., 2010). Atomic force

microscopy has also revealed the existence of dimers of homotrimers between P2X4 receptors and either P2X2 or P2X7 receptors, suggesting oligomeric interactions between P2X receptors (Antonio et al., 2011). However, as stated above, we found no evidence for specific BRET or BiFC between P2X2 and P2X4 subunits, supporting that the BRET observed between P2X subunits results exclusively from heterotrimeric assembly rather than from oligomeric interactions. This is further supported by our cross-linking experiments with BS3, which clearly demonstrate that P2X2 and P2X5 subunits only associate within a trimeric complex.

Trimeric P2X2/5 receptors present two possible stoichiometries, P2X2/(P2X5)₂ and (P2X2)₂/P2X5. These findings were established by measuring the energy transfer between one given luciferase-tagged subunit and two hemi-YFP tagged subunits of the second kind. This dual stoichiometry was also confirmed with a biochemical approach, by coexpressing Myc- and Cam-tagged subunits, although in these experiments, only one cameleon-

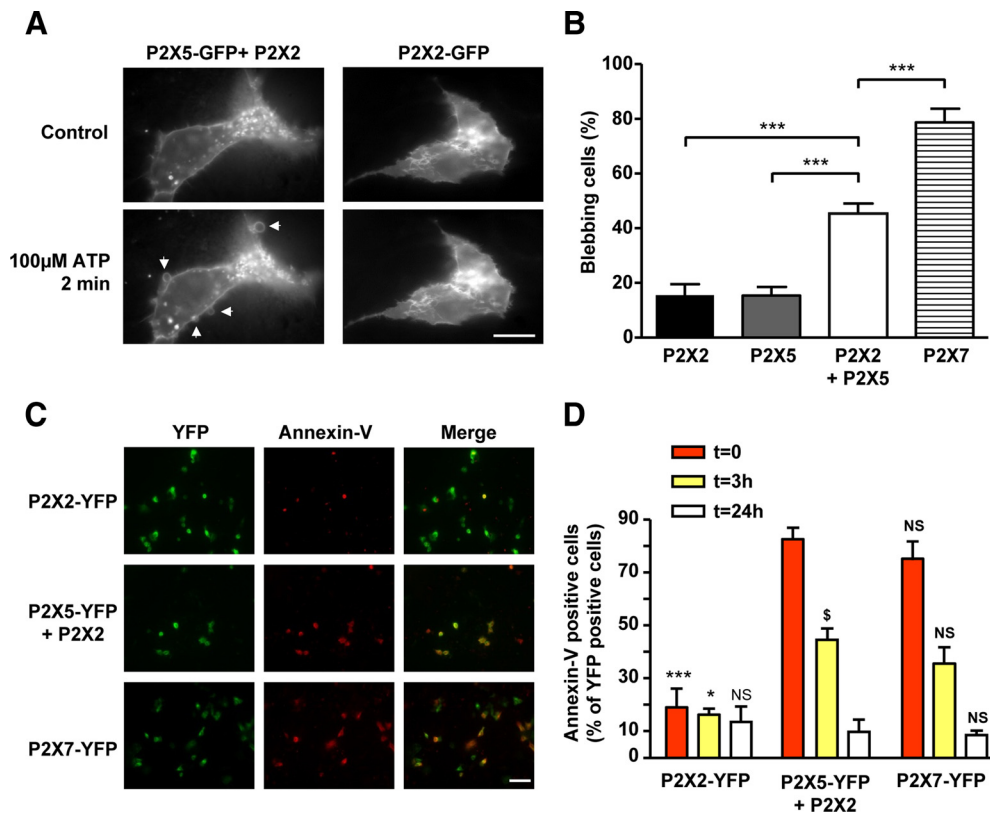


Figure 9. Activation of P2X2/5 receptors mediates membrane blebbing and pseudo-apoptosis. **A, B**, P2X2/5 activation triggers membrane blebbing. **A**, Representative fluorescence images of HEK cells expressing P2X2-GFP or P2X2 + P2X5-GFP 2 min after 100 μ M ATP stimulation. Arrows indicate ATP induced blebs. Scale bar, 10 μ m. **B**, Quantitative analysis of the percentage of blebbing cells expressing P2X2-GFP, P2X5-GFP, P2X2/5-GFP, or P2X7-GFP after stimulation with 100 μ M ATP (2 mM for P2X7) for 2 min. Results were normalized to the number of GFP-positive cells. Data are mean \pm SEM of $N = 3$ independent experiments. $***p < 0.005$, one-way ANOVA, followed by Bonferroni's multiple-comparison test. **C, D**, P2X2/5 receptor activation induces transient annexin-V exposure. **C**, HEK cells transfected with P2X2-YFP, P2X2 + P2X5-YFP, or P2X7-YFP were stimulated with 100 μ M ATP (500 μ M for P2X7-YFP) and annexin-V staining was performed 5 min after the end of the stimulation. Scale bar, 50 μ m. **D**, Analysis of annexin-V staining at 0, 3, or 24 h after ATP stimulation. Experiment was performed as in **C**, except that annexin-V staining was performed at the time indicated after ATP stimulation. In all panels, results were normalized to the number of YFP-positive cells. Data are mean \pm SEM, $n > 30$ cells, $N = 3$ experiments. $*p < 0.05$, $***p < 0.005$; N.S., not significant by comparison with P2X2/5 transfected cells at the same time point. $^{\$}p < 0.05$, comparison between P2X2/5-expressing cells at 0 and 3 h. One-way ANOVA, followed by Bonferroni's multiple-comparison test.

tagged subunit was incorporated into a receptor complex. However, this was equally observed for homomeric P2X5 and P2X2, as well as heteromeric subunit assemblies. It is likely that, because of the size of the fusion partner, Cam-tagged subunits have either reduced expression efficiency or slower maturation, resulting in a lower efficacy of incorporation within trimers. Alternatively, channels associating two Cam-tagged subunits and one Myc could be present but remain undetected due to the low sensitivity of the anti-Myc antibody and the presence of a single Myc-tagged subunit within the complex. Dual stoichiometry has been reported for the assembly of P2X2 and P2X6 subunits, for which stoichiometry is influenced by the relative amount of each subunit (Barrera et al., 2007). Alternate stoichiometries have also previously been demonstrated for the pentameric assembly of nicotinic $\alpha 2\beta 4$ channels (Moroni et al., 2008; Carbone et al., 2009). Whether these alternate assemblies of ligand-gated channels result from the heterologous expression system used, or are of physiological relevance, remains to be established.

P2X5 orthologues display considerable functional diversity. Bullfrog, chicken, and human homomeric P2X5 receptors show robust expression in oocytes or mammalian cells (Jensik et al., 2001; Ruppelt et al., 2001; Bo et al., 2003) and display specific properties such as high chloride permeability, or, in the case of the full-length human isoform, a quasi-instantaneous NMDG permeability. This is in stark contrast to rodent homomeric P2X5

receptors, which are poorly functional (Collo et al., 1996; Cox et al., 2001). Our functional data, together with previous studies on P2X1/5 receptors (Haines et al., 1999; Lê et al., 1999; Surprenant et al., 2000), support the idea that rodent P2X5 subunits are auxiliary subunits, associating with P2X1, P2X2, and P2X4 subunits to generate molecular diversity among P2X receptors. Rat P2X2 and P2X5 receptors, however, have similar pharmacology (North and Surprenant, 2000). Consistent with this, our pharmacological profiling shows that all of the agonists tested had similar efficacies at P2X2 and P2X2/5 receptors, although the potencies were significantly lower at P2X2/5 receptors. Only BzATP displayed a much lower efficacy at P2X2/5 receptors, which may be due to the lack of sensitivity of the rat homomeric P2X5 receptor to BzATP (North and Surprenant, 2000). Rescue experiments using a trafficking mutant of either P2X2 or P2X5 subunits did not allow the unambiguous definition of the pharmacological profile of P2X2/5 heteromers. These experiments, however, suggest that (P2X2)₂/P2X5 might be the only functional heteromer, since (1) P2X5 subunits are unable to functionally rescue P2X2-K366A and (2) upon coexpression of P2X2 and P2X5-K372A subunits, BzATP potency is significantly lower than at P2X2. However, in this latter experiment, the presence of homomeric P2X2 receptors prevented any precise pharmacological profiling.

Our immunohistochemical data are consistent with the expression of P2X2/5 receptors in proprioceptive neurons of the

mesencephalic trigeminal nucleus. Indeed, these neurons express P2X5 receptors (Collo et al., 1996; Cook et al., 1997), as well as P2X receptors with a pharmacology differing from that of any recombinant homomeric or heteromeric receptors (Cook et al., 1997; Patel et al., 2001). Similarly, in sensory neurons, P2X2/5 receptors could contribute to the recently described P2X-mediated ASIC3 facilitation involved in the detection of muscle ischemia (Birdsong et al., 2010).

Similar to P2X7 receptors, P2X2 and P2X4 receptors can undergo time-dependent permeability changes, leading to significant permeability to NMDG (Khakh et al., 1999; Virginio et al., 1999b). P2X2/5- and P2X7-evoked conductances have some striking similarities. In oocytes, compared with P2X2, the P2X2/5-evoked NMDG conductance developed with a delayed onset and slower kinetics, was not saturable and poorly reversible, and displayed a complete loss of the initial inward rectification. P2X2/5-evoked sustained NMDG conductances were also observed in HEK cells, ruling out a cell-type dependent phenomenon. In addition, as described for P2X7 (Jiang et al., 2005), the P2X2/5 sustained NMDG conductance closes upon switching to sodium-containing extracellular solution. In HEK cells, P2X2/5 receptor-mediated Yo-Pro-1 uptake displayed slower kinetics as well as lower sensitivity to ATP, compared with P2X2 receptors. Nevertheless, the absolute value of fluorescence was larger in cells transfected with P2X2/5. Additionally, under physiological conditions, P2X2/5- and P2X7-evoked EtBr uptake shared common features not observed with P2X2, mainly slow kinetics and an absence of saturation. Altogether, these data support the notion that activation of P2X2/5 receptors triggers membrane permeabilization, a function thought to be uniquely associated with P2X7 receptors (Skaper et al., 2010). It remains to be determined whether NMDG and Yo-Pro-1 uptake are mediated by the same molecular mechanisms at P2X2 and P2X2/5 receptors. Our observation that the conformational motions of the cytoplasmic tail of P2X2/5 receptor are undetectable by BRET might support two independent mechanisms.

P2X2/5 receptors share other functional properties with the P2X7 receptor, particularly its ability to trigger membrane blebbing (Morelli et al., 2003; Verhoef et al., 2003) and pseudoapoptosis (Virginio et al., 1999a; Mackenzie et al., 2005). Blebs are often associated with apoptosis; however, in the case of P2X7 receptors, short ATP stimulations trigger blebbing and plasma membrane PS flip that do not result in cell death (Mackenzie et al., 2005), a process qualified as pseudoapoptosis. P2X2/5 receptor-induced blebbing displays similar features to P2X7. It is a reversible process observed under physiological conditions with an onset of <20 s, and is highly dynamic with blebs completely resorbing while new ones appear. Furthermore, activation of P2X2/5 receptors also results in pseudoapoptosis, characterized by a reversible annexin-V exposure associated with an absence of cell death.

ATP-evoked Yo-Pro-1 uptake, blebbing, or apoptosis are widely considered to be specific signatures of P2X7 receptor activity. Our data establish that under physiological conditions, identical phenotypes can also result from activation of P2X2/5 heteromeric receptors. In the nervous system and particularly in neurons, the expression of P2X7 receptors is somewhat controversial (Sim et al., 2004; Anderson and Nedergaard, 2006). It is thus conceivable that in these structures or cells, other P2X receptors might contribute to cellular responses typically attributed to P2X7 receptors.

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