Peripheral Inflammation Sensitizes P2X Receptor-Mediated Responses in Rat Dorsal Root Ganglion Neurons

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ATP-gated P2X receptors in nociceptive sensory neurons participate in transmission of pain signals from the periphery to the spinal cord. To determine the role of P2X receptors under injurious conditions, we examined ATP-evoked responses in dorsal root ganglion (DRG) neurons isolated from rats with peripheral inflammation, induced by injection of complete Freund’s adjuvant (CFA) into the hindpaw. Application of ATP induced both fast- and slow-inactivating currents in control and inflamed neurons. CFA treatment had no effect on the affinity of ATP for its receptors or receptor phenotypes. On the other hand, inflammation caused a twofold to threefold increase in both ATP-activated currents, altered the voltage dependence of P2X receptors, and enhanced the expression of P2X2 and P2X3 receptors. The increase in ATP responses gave rise to large depolarizations that exceeded the threshold of action potentials in inflamed DRG neurons. Thus, P2X receptor upregulation could account for neuronal hypersensitivity and contribute to abnormal pain responses associated with inflammatory injuries. These results suggest that P2X receptors are useful targets for inflammatory pain therapy.

Key words: dorsal root ganglion; ATP; P2X receptor; Western blotting; peripheral inflammation; pain; electrophysiology

In addition to being an intracellular energy source, ATP is released from neuronal and non-neuronal cells, acts on purinergic receptors, and regulates activities of autonomic and sensory neurons, smooth muscle, and endothelial cells (Jahr and Jessell, 1983; Cook and McCleskey, 2000; North and Surprenant, 2000; Burnstock, 2001). In primary sensory dorsal root ganglion (DRG) cells, ATP plays a prominent role in signaling. It depolarizes DRG neurons by eliciting fast- and slow-inactivating inward currents. The fast-inactivating ATP currents are mediated by homomeric P2X3 receptors; the slow-desensitizing currents are mediated by heteromeric P2X2/3 receptors (Bradbury et al., 1998; Virginio et al., 1998; Burgard et al., 1999; Grubb and Evans, 1999). In addition, ATP modulates synaptic transmission at DRG and dorsal horn synapses (Bardoni et al., 1997; Li et al., 1998). It enhances spontaneous glutamate responses and elicits action potentials to evoke glutamate release (Gu and MacDermott, 1997; Nakatsuka and Gu, 2001). Studies of P2X receptor distributions show that P2X2 and P2X3 receptor subtypes are selectively expressed in structures associated with pain signal processing, including small- and medium-sized DRG neurons, peripheral and central sensory terminals, and superficial dorsal horns (Chen et al., 1995; Lewis et al., 1995; Cook et al., 1997; Vulchanova et al., 1998; Kanjhan et al., 1999). Behavioral experiments suggest that applications of the P2X agonists ATP and αβmeATP to the rat hindpaw decrease the tail-flick latency and produce flinching and writhing behaviors (Cockayne et al., 2000; Souslova et al., 2000; Tsuda et al., 2000). Thus, activation of P2X receptors in sensory neurons facilitates transmission of nociceptive signals from the periphery to the spinal cord.

The consequences of nerve and tissue injuries on ATP responses have not been thoroughly explored. Insults to afferent fibers and peripheral tissues, such as neuropathy and inflammation, frequently give rise to exaggerated responses to non-noxious and noxious stimuli (alldynia and hyperalgesia). These pathological responses are thought to arise from sensitization of DRG and dorsal horn neurons to external stimuli (Woolf and Doubell, 1994; Xu et al., 2000). Stanfa et al. (2000) find that the spinally administered P2X antagonists suramin and pyridoxal phosphate-6-azophenyl-2,4'-disulfonic acid (PPADS) reduce C-fiber-evoked discharges in deep dorsal horn neurons of rats with inflammation but have no effect on those in normal or nerves-intact rats. Hamilton et al. (1999) show that high concentrations of ATP (≥100 nmol) to the hindpaw of normal rats are required to produce nocifensive behaviors (i.e., paw lifting, shaking, and licking) and heat hyperalgesia. However, 1 nmol of ATP can evoke these behaviors in rats inflamed with carrageenan. Furthermore, ATP-evoked activity of C-mechanoreceptor or polymodal nociceptors is greatly enhanced (Hamilton et al., 2001). P2X receptor activation is therefore facilitated after inflammation. The mechanism underlying the facilitation is unknown. Here we examine the ATP-evoked responses and the expression of P2X receptors in DRG neurons isolated from rats with peripheral inflammation. Our results show that inflammation produces a large increase in P2X receptor currents.

Parts of this work have been published previously in abstract form (Xu and Huang, 1999).

MATERIALS AND METHODS

Induction of peripheral inflammation. All experiments were approved by the Institutional Animal Care and Use Committee at the University of Texas Medical Branch and were in accordance with the guidelines of the National Institutes of Health and the International Association for the Study of Pain. Sprague Dawley rats (27–37 d old) were used for the
dissolved in 40 mM NaCl, 5KCl, 2.0 K$_2$HPO$_4$, 1.5 CaCl$_2$, 6 MgCl$_2$, 10 glucose, and 10 HEPES, pH 7.2 (osmolarity, 305 mOsm). After removal of the connective tissue, the ganglia were transferred to a 10 ml dissecting solution containing collagen IV (1.0–1.5 mg/ml; Boehringer Mannheim, Indianapolis, IN) and trypsin (1.0 mg/ml; Sigma, St. Louis, MO) and incubated for 1 hr at 34.5°C. DRGs were then taken from the enzyme solution, washed, and put in 3 ml of the dissecting solution containing DNase (0.5 mg/ml, Sigma). Cells were subsequently dissociated by trituration with fire-polished glass pipettes and placed on acid-cleaned glass coverslips.

**Perforated patch recording and application of drugs.** Cells were superfused (2 ml/min) at room temperature with an external solution containing (in mM): 130 NaCl, 5 KCl, 2.0 K$_2$HPO$_4$, 2.5 CaCl$_2$, 1.5 MgCl$_2$, 10 HEPES, and 10 glucose, pH 7.2 (osmolarity, 295–300 mOsm). ATP-induced currents and tension potentiation were recorded using the perforated patch-clamp technique. The patch electrode had a resistance between 2.2 and 3.5 MΩ. The pipette tip was initially filled with amphotericin-free pipette solution, containing (in mM): 100 K$_2$SO$_4$, 40 KCl, and 10 HEPES, pH 7.25 adjusted with KOH (osmolarity, 290 mOsm). The pipette was then backfilled with the same pipette solution containing amphotericin B (300 μg/ml). The currents were filtered at 2–5 kHz and sampled at 50 or 100 μsec per point.

All experiments were performed within 2 h of the animals being killed. Control and CFA-treated rats were examined under voltage-clamp conditions. Peripheral inflammation was induced by injecting CFA into the ankle and plantar surface of the rat left hindpaw. L4–L6 DRGs were isolated 3–14 d after the CFA injection, a period of peak hyperalgesic conditions. We chose small to medium diameter (15–40 μm) DRG neurons for the study because they mediate transmission of nociceptive signals (Willis and Coggeshall, 1991).

Applications of ATP (20 μM) evoked large inward currents at ~60 mV holding potential in control DRG neurons. Based on the time course of the responses, they were categorized as fast, slow, and mixed responses. The fast ATP-evoked currents were rapidly activating and inactivating (Fig. 1A). The rise times of the fast responses were short ($T_{\text{r}}$ = 6.5 ± 1.1 msec; $n$ = 18). The currents reduced to <5.0 ± 0.5% of their peak amplitudes within 2 sec of ATP applications. The inactivating phase of the currents was best fitted with a sum of two exponentials ($\tau_{\text{in}}$ = 38.1 ± 7.7 msec, $A_1$ = 605.3 ± 171.2 pA, $\tau_{\text{in}}$ = 477.3 ± 177.2 msec, $A_2$ = 120.2 ± 20.6 pA; $n$ = 11). Once inactivated, the fast ATP responses recovered very slowly. The original peak amplitude would not be restored unless there was a 10–15 min wait between consecutive ATP applications.

The slow ATP responses in control neurons were characterized by relatively slow rise times ($T_{\text{r}}$ = 38.5 ± 5.97 msec; $n$ = 10). The inactivation kinetics varied among cells. In a small percentage (~8%) of cells, slow ATP responses showed very little inactivation during the ATP application. A majority of slow responses, however, showed inactivation. The decay kinetics could be fit with one exponential. The average inactivation time constant ($\tau_{\text{in}}$) was 207.5 ± 200.4 msec ($n$ = 10), which was ~4–50 times slower than those of the fast ATP responses. Unlike the fast ATP responses, the slow ATP responses recovered rather quickly. The peak response returned to its original size within 1 min after an ATP application.

ATP also evoked inward currents in a large number of neurons isolated from CFA rats. Similar to control cells, ATP evoked fast and slow responses in inflamed neurons. The most prominent change was the large enhancement of current amplitudes (Fig. 1B). The average peak current density of the fast responses in inflamed neurons was 2.7-fold larger (control, 0.30 ± 0.05 pA/μm$^2$, $n$ = 29; CFA, 0.82 ± 0.13 pA/μm$^2$, $n$ = 48). The kinetic characteristics of the fast ATP responses of inflamed neurons were similar to those obtained from control neurons (Table 1).
The fast ATP responses in inflamed neurons activated rapidly ($T_{\text{a}} = 6.3 \pm 0.78$ msec, $n = 19$) and desensitized in two phases ($\tau_{\text{on}} = 49.4 \pm 7.9$ msec, $A_1 = 1352.7 \pm 237.1$ pA; $\tau_{\text{off}} = 292.7 \pm 30.4$ msec, $A_2 = 325.5 \pm 58.4$ pA; $n = 19$). The currents were desensitized completely during the 2 sec ATP application.

ATP also evoked slow responses in inflamed neurons. The amplitudes of slow responses were also greatly potentiated. The average peak current density of the slow-inactivating currents was increased by 2.8-fold (control, peak $0.24 \pm 0.05$ pA/μm², $n = 25$; CFA, peak $0.68 \pm 0.09$ pA/μm², $n = 38$); the average steady-state slow current density of the slow responses was increased by 3.0-fold (control, $0.10 \pm 0.02$ pA/μm², $n = 25$; CFA, $0.30 \pm 0.05$ pA/μm², $n = 38$) (Fig. 1C). The kinetic properties of slow responses in inflamed neurons were not significantly different from those of control neurons (Table 1). The slow ATP responses of inflamed neurons had a mean rise time ($T_{\text{a}}$) of $52.0 \pm 8.2$ msec ($n = 16$), and a mean decayed time constant of $\tau_{\text{on}} = 2318.4 \pm 379.5$ msec ($n = 16$).

We also observed ATP currents with mixed fast and slow characteristics in both control and inflamed (data not shown). Like the fast ATP responses, the mixed ATP responses were characterized by a fast rise time and a distinct two-phase inactivation. The fast inactivation phase had a time constant similar to the $\tau_{\text{on}}$ of the fast ATP responses. However, the slow inactivation phase had a time constant that was much slower than the $\tau_{\text{off}}$ of the fast ATP responses. Thus, a substantial portion of the current remained at the end of the 2 sec ATP application, a characteristic feature of the slow ATP responses. To simplify our analyses, the mixed ATP responses, which occurred in 8.5% of ATP responding control neurons (12 of 141 cells tested) and 11.6% of responding CFA neurons (32 of 276 cells), were not included in this study.

**Cell distribution of ATP responses**

We then analyzed the cell types that displayed either fast or slow ATP responses. ATP-induced responses were observed in 89.4% of all recorded DRG neurons ($n = 141$) isolated from control rats and in 93.8% of DRG neurons ($n = 276$) isolated from CFA-injected rats. Thus, the percentages of neurons responding to ATP remained unchanged after CFA treatment ($p > 0.05$). Analyses of the types of ATP responses in control neurons indicated that 33.3% of recorded neurons ($n = 47$) exhibited fast-inactivating ATP currents, and 47.5% of cells exhibited slow-inactivating ATP currents ($n = 67$). After inflammation, 42.8% of neurons ($n = 118$) exhibited fast ATP responses, and 39.5% of cells ($n = 109$) exhibited slow ATP responses. Therefore, two types of responses occurred with approximately equal frequencies in control and CFA-treated rats ($p > 0.05$) (Fig. 2A).

Cell size distribution of fast- and slow-inactivating ATP responses was also obtained using cumulative distribution analyses. The percentages of cells exhibiting ATP responses versus cell diameters smaller than the indicated values were plotted (Fig. 2B). We found that 50% of cells responding to ATP with fast-inactivating currents had diameters <26 μm in both control and CFA rat groups. Cells responding to ATP with slow-inactivating currents for both rat groups were significantly larger (i.e., 50% of cells responding with slow currents had diameters <33 μm in control rats and <31 μm in CFA rats). Because the cell size distributions for both ATP responses are the same for normal and inflamed rats, inflammation does not appear to alter the types of DRG cells expressed P2X receptors.

**P2X receptor phenotypes**

ATP activates more than one subtype of P2X receptors in control DRG neurons (Vulchanova et al., 1997; North and Surprenant, 2000). It is of interest to determine whether the same P2X receptor subtypes are expressed in DRGs after inflammation. Antagonists were first used to identify P2X receptors in DRGs. Suramin (30 μM) and PPADS (50 μM) completely blocked fast and slow ATP-evoked currents in control ($n = 20$) and inflamed ($n = 25$) neurons (Fig. 3A, left and middle). These two antagonists, at tens of micromolar concentrations, are known to block homomeric P2X1, P2X2, P2X3, and P2X5 and heteromeric P2X2/3 receptors without significantly affecting homomeric P2X4, P2X6, and P2X7 receptors (North and Barnard, 1997; North and Surprenant, 2000). Thus, P2X4, P2X6, and P2X7 were not present in either control or inflamed DRGs. We then used the antagonist TNP-ATP to determine whether homomeric P2X2 receptors were present in our DRG neurons (Fig. 3A, right).
Table 1. Kinetics of ATP-induced fast and slow currents in dorsal root ganglion neurons isolated from control (CON) and CFA-treated rats

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<td>CON</td>
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<td>CFA</td>
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Data represent the mean ± SEM; \(n\) represents the cell number. The differences in \(\tau_m\) and \(\tau_m\) values between CON and CFA rat groups were not significant (\(p > 0.05\)).

Figure 2. Inflammation does not change the percentages and the size distributions of neurons responding to ATP. A. Percentages of responding cells. The percentage of the total number of cells (Total) responding to ATP was 89.4% (\(n = 141\)) in control rats (CON) and 93.8% (\(n = 276\)) in CFA rats. The change was not significant (\(\chi^2\) test; \(p > 0.05\)). The percentages of cells with fast-inactivating ATP responses (Fast) (control, 33.3%, \(n = 47\); CFA, 42.8%, \(n = 118\)) and the percentages of cells with slow-inactivating ATP responses (Slow) (control, 47.5%, \(n = 67\); CFA, 39.5%, \(n = 109\)) were not altered by inflammation. B. Cell size distributions for ATP responses. Distribution of cell diameter was expressed in cumulative histograms, i.e., percentages of cells that responded with either fast or slow ATP responses versus cell diameters that were smaller than the indicated values. In control rats, 50% of cells responding to ATP with fast-inactivating currents (13 of 27 cells tested) had diameters 26 \(\mu\)m; 50% of the cells responding to ATP with slow-inactivating currents (23 of 46 cells tested) had diameters <33 \(\mu\)m. The size difference was significant (\(p < 0.05\); Kolmogorov–Smirnov test). In CFA rats, 50% of the cells responding to ATP with fast-inactivating currents had diameters <26 \(\mu\)m (32 of 65 cells tested); 50% of the cells responding to ATP with slow-inactivating currents had diameters <31 \(\mu\)m (28 of 56 cells tested). CFA treatment did not change the size distribution of cells responding to ATP with either the fast- or slow-inactivating currents.

TNP-ATP is 500-fold more sensitive to homomeric P2X1 and P2X3 and heteromeric P2X2/3 receptors than to homomeric P2X2 receptors (Thomas et al., 1998). High concentrations (\(\geq 1 \muM\)) of TNP-ATP should block P2X1, P2X3, and P2X2/3 receptor-mediated responses but leaves most homomeric P2X2 receptor-mediated responses intact (Thomas et al., 1998; Virginio et al., 1998; North and Surprenant, 2000). At 1 \(\muM\), TNP-ATP blocked all fast ATP currents and reduced slow ATP currents by 98% in control (\(n = 18\)) and inflamed (\(n = 24\)) neurons (Fig. 3A, right). Thus, the responses mediated by homomeric P2X2 receptors, if present in our cells, would be small.

We then used the P2X receptor agonist \(\alpha\beta\text{meATP}\) to further identify P2X receptor types in control and inflamed neurons. Unlike ATP, \(\alpha\beta\text{meATP}\) has low affinity for homomeric P2X2 and P2X5 receptors (North and Surprenant, 2000). If P2X2 and/or P2X5 receptors were present in significant quantities, a saturated concentration of ATP (100 \(\muM\)) or \(\alpha\beta\text{meATP}\) (100 \(\muM\)) should evoke different responses. This was not observed. ATP and \(\alpha\beta\text{meATP}\) activated currents of similar amplitudes in control (\(n = 9\)) and inflamed (\(n = 12\)) neurons (Fig. 3B). Homomeric P2X2 and P2X5 receptors, therefore, were not present in sufficient amount to contribute to ATP responses in either control or inflamed DRGs. The expression of P2X1 receptor in control and inflamed neurons has not been studied in detail. Preliminary Western blot analyses showed that the P2X1 receptor immunoreactivity was low in both control and inflamed neurons, suggesting that P2X1 was not the major receptor type in DRGs (data not shown). From these experiments, we conclude that homomeric P2X3 and heteromeric P2X2/3 receptors are the main receptor types in inflamed DRGs and that inflammation does not elicit significant changes in P2X receptor phenotypes.

Affinity of ATP for P2X receptors

To determine whether the increase in ATP responses in inflamed neurons arises from changes in the affinity of ATP for P2X receptors, dose–response curves for ATP in control and inflamed rat groups were studied (Fig. 4). ATP, at \(\geq 100 \muM\), elicited both maximal fast and slow ATP responses. The maximal fast response was 2.5-fold larger and the maximal slow response was 2.3-fold larger in inflamed neurons. Dose–response curves for both fast and slow ATP responses were fit with the Hill equation. The \(E_{50}\) for fast ATP responses was 1.7 ± 0.9 \(\muM\) in control and 2.0 ± 0.69 \(\muM\) in inflamed neurons (Fig. 4A). The \(E_{50}\) for slow ATP responses was 5.7 ± 1.4 \(\muM\) in control and 3.6 ± 1.2 \(\muM\) in inflamed neurons (Fig. 4B). The changes in ATP affinities for P2X receptors in inflamed DRG neurons were not significant.

Leftward shift of conductance–voltage curves

The voltage dependence of ATP responses was also determined. Currents in response to ATP applications were measured at different holding potentials. The peak currents versus voltage (\(I–V\)) curves were plotted. Both fast and slow ATP currents reversed at near +10 mV in control cells (Fig. 5A), and CFA treatment did not change the reversal potentials of ATP re-
sponses (Fig. 5). Therefore, inflammation had no significant effect on the permeation properties of P2X3 and P2X2/3 receptors.

Both fast- and slow-inactivating ATP currents showed inward rectification (Fig. 5). The conductance–voltage ($G-V$) relationships of both types of currents were fit with the Boltzmann equation (Fig. 6). The $G-V$ curves obtained for the fast ATP responses in control neurons had a $Z = 0.97 \pm 0.07$, $G_{\text{max}} = 4.4 \pm 0.7 \text{pS/\mu m}^2$, and $V_{0.5} = -35.6 \pm 2.9 \text{mV}$ ($n = 10$). CFA treatment did not significantly change the $Z$ ($0.91 \pm 0.08; n = 12$) of the $G-V$ curve. As expected from the current data (Fig. 1), the $G_{\text{max}}$ of inflamed neurons was 2.9-fold larger, i.e., $G_{\text{max}} = 12.9 \pm 1.4 \text{pS/\mu m}^2$ ($n = 12$). Furthermore, the $V_{0.5}$ shifted significantly in the hyperpolarized direction (Fig. 6) ($V_{0.5} = -49.5 \pm 2.3 \text{mV}; n = 12$).

The $G-V$ of the slow-inactivating ATP currents in control neurons had $Z = 0.92 \pm 0.09$, $G_{\text{max}} = 5.0 \pm 0.9 \text{pS/\mu m}^2$, and $V_{0.5} = -25.0 \pm 4.8 \text{mV}$ ($n = 6$). Compared with the fast ATP currents, the slow ATP responses inactivated at a more depolarized potential. Inflammation did not affect $Z$ ($0.95 \pm 0.06; n = 12$), but it increased the $G_{\text{max}} (11.0 \pm 1.4 \text{pS/\mu m}^2; n = 12)$ of slow ATP responses by 2.2-fold and shifted $G-V$ curves in the hyperpolarized direction ($V_{0.5} = -43.5 \pm 2.8 \text{mV}; n = 12$). Thus, in addition to increasing the maximal conductances, inflammation causes both types of ATP responses to inactivate at more hyperpolarized potentials.

### Increased membrane depolarization

We then compared the effect of ATP on the membrane depolarization of DRG neurons in control and CFA neurons under current-clamp conditions. The average resting membrane potential of DRG neurons recorded from CFA-treated rats was $-49.9 \pm 0.7 \text{mV}$ ($n = 104$), which was not significantly different from the resting membrane potential of neurons recorded from control rats ($-50.8 \pm 1.2 \text{mV}; n = 54$). In control rats, application of ATP (20 \text{mM}) produced depolarizations of membrane potentials in 18 of 22 cells tested (Fig. 7A). Most of the depolarizations were subthreshold (Fig. 7A,B, top left). After CFA treatment, ATP induced depolarization in 30 of 34 neurons. All of the depolarizations were large enough to evoke action potentials.
Figure 5. ATP currents in control and inflamed neurons exhibit steep voltage dependence. Examples of current–voltage (I–V) relationships of peak fast (left) and slow (right) ATP currents in control (CON) (A) and CFA (B) neurons. The currents were measured at different holding potentials. The current traces were shown in the inset of each I–V curve. The reversal potentials of the currents did not change after CFA treatment. Data were obtained from four different cells.

(Fig. 7A,B, bottom right). Because Na⁺ channels are upregulated in the inflammatory state (Gould et al., 1998; Gold, 1999) and could affect changes in the firing properties of inflamed neurons, their contribution to the depolarization has to be eliminated. We therefore isolated the depolarization attributable to P2X receptor activation by using TTX to block TTX-sensitive Na⁺ channels and a depolarized prepulse to inactivate both TTX-sensitive and -resistant Na⁺ channels (Ogata and Tatebayashi, 1993; Rush et al., 1998). TTX (2 μM) could block cell firings in ~50% of the DRG cells isolated from control and CFA rats. In the other 50% of the cells tested, TTX had little effect on the spike generation. The ATP-evoked depolarizations in TTX-sensitive and -resistant neurons were evaluated separately. In TTX-sensitive neurons isolated from control rats (Fig. 7A, top), the average size of the depolarization was 12.4 ± 3.7 mV (n = 4) before TTX and 11.8 ± 4.1 mV (n = 4) after TTX. To inactivate TTX-resistant Na⁺ channels that might also be present in these cells, an 8 sec depolarized prepulse to −10 or −15 mV was applied before ATP application. ATP-evoked depolarization, after the prepulse, was 11.3 ± 3.6 mV (n = 4). The sizes of depolarizations under the various experimental conditions were not significantly different. We then examined ATP-evoked depolarizations in neurons isolated from inflamed rats. ATP evoked cell firings in all of the TTX-sensitive inflamed neurons (Fig. 7A, bottom). The depolarization evoked by ATP was 30.9 ± 1.4 (n = 6) with TTX and 30.8 ± 1.5 mV (n = 6) with both TTX and the prepulse. Thus, ATP evoked a substantially larger depolarization after inflammation (Fig. 7B).

The same experiments were repeated in TTX-resistant neurons. ATP did not evoke cell firings in most TTX-resistant neurons isolated from control rats (Fig. 7C, top). The average ATP-evoked depolarization in these cells was 15.7 ± 1.6 mV (n = 4) before TTX and 15.9 ± 1.7 mV (n = 4) after TTX. When a depolarized prepulse was applied to this cell group, action potentials were often evoked at the beginning of the prepulse, even in the presence of TTX. The firing then subsided as the TTX-resistant Na⁺ channels became inactivated during the prepulse. The depolarization evoked by ATP applied after the prepulse was 15.6 ± 1.7 mV (n = 4). The contribution of Na⁺ channel activation to ATP depolarization was not significant. ATP invariably evoked cell firing in responsive TTX-resistant neurons isolated from CFA-treated rats (Fig. 7B, bottom traces). These spikes could not be blocked by TTX but were inactivated by the depolarized prepulse. The average ATP-induced depolarization after the depolarized prepulse was 32.4 ± 1.5 mV (n = 4), which again is much larger than that obtained in control neurons (Fig. 7D).

We then examined threshold voltages of action potentials in...
both control and inflamed neurons. The threshold voltage was $-24.24mV (n=9)$ in control neurons and $-25.53mV (n=25)$ in inflamed neurons. CFA treatment did not change the threshold voltage significantly. Thus, in both TTX-sensitive and -resistant neurons isolated from control rats, TTX-evoked depolarizations are subthreshold. In contrast, ATP evoked depolarizations in both types of neurons isolated from CFA rats are large and exceed the firing threshold of the neurons.

**Enhanced P2X receptor expression**

To determine whether the expression of P2X receptors indeed increases in DRG after inflammation, Western blotting assays were performed on DRGs in control rats and in inflamed rats ipsilateral to the CFA-injected paw. Proteins were isolated from L4–L6 DRGs of control rats and rats treated with CFA for 5 d. After separating the proteins by electrophoresis under denaturing conditions, they were transferred to nylon membranes and probed with anti-P2X2 and anti-P2X3. Anti-P2X2 antibody labeled a $~64$ kDa molecular weight protein, and anti-P2X3 labeled a $~57$ kDa protein. After CFA treatment, the molecular weight of the proteins did not change. However, the level of expression of both P2X2 and P2X3 receptors was increased significantly (Fig. 8) (P2X2, CFA/control = 1.81; P2X3, CFA/control = 1.82). Thus, inflammation upregulates the P2X2 and P2X3 receptor expression in DRGs.

**DISCUSSION**

We show here that ATP responses in DRG neurons are altered by inflammation. The most prominent change is a twofold to threefold increase in the current density of both fast and slow ATP responses (Fig. 1). Because the EC$_{50}$ of the dose–response curve for ATP does not change in inflamed neurons (Fig. 4), the increase cannot be attributed to an increase in the affinity of ATP for its receptors. Possible mechanisms for the potentiation of ATP currents include an increase in single-channel conductance, enhancement of channel opening probability, and/or upregulation of P2X receptor expression. Although single ATP receptor channel properties in inflamed neurons have yet to be studied, an increase in the opening probability of ATP channels is not likely because the kinetic properties of both fast and slow ATP currents remain unchanged after CFA treatment (Table 1). Because CFA produces a significant increase in P2X2 and P2X3 proteins (Fig. 8), upregulation of P2X receptor expression is a major cause for
The large increase in ATP responses after the development of inflammation.

The current-voltage curves of both slow and fast ATP-evoked currents show inward rectification in control and CFA-treated rats (Fig. 5). The steepness of the rectification is similar to those reported by others in normal rats (Krishtal et al., 1983; Bean, 1990). The inward rectification characteristics of ATP responses are thought to arise from voltage-dependent blocking of intracellular cations (Krishtal et al., 1988) and/or fast voltage-dependent gating of ATP-activated channels (Bean, 1990; Bean et al., 1990). Our analyses of the voltage dependence of ATP responses show that the G-V curves of both fast and slow ATP responses shift in the hyperpolarized direction after inflammation (Fig. 6). The mean \( V_{\text{p,50}} \) of fast ATP current shifts from \(-35.6 \) to \(-49.5 \) mV; the mean \( V_{\text{p,50}} \) of slow responses shifts from \(-25.4 \) to \(-43.5 \) mV. The mechanism underlying the shift is yet unclear. Changes in the phosphorylating state of P2X receptors after inflammation could be a contributing factor (Paukert et al., 2001). One physiological consequence of the inward rectification of ATP currents is regulation of action potential generation. A large inward ATP current generated below the firing threshold (less than \(-25 \) mV) will depolarize cells quickly and thus activate voltage-dependent ion channels. As the membrane potential depolarizes, the inward ATP current will get smaller. When the membrane potential becomes positive, the outward ATP current is nearly blocked.

Thus, activation of P2X receptors will facilitate the generation of action potential without shunting it at positive potentials. When the G-V curves of ATP responses shift to hyperpolarized potentials after CFA treatment, the depolarizing effects of P2X receptor activation would be dampened because a smaller fraction of P2X receptors are activated at the resting potential. In our case, the relative conductance \((G/G_{\text{max}})\) of the fast ATP response at \(-50 \) mV is 0.59 in control cells but becomes 0.51 in CFA neurons. A 2.9-fold increase in \( G_{\text{max}} \) after the development of inflammation would give a 2.5-fold increase in conductance at \(-50 \) mV. The steepness of the G-V curve, the increase in conductance in inflamed neurons is still large.

We also compared the ATP-induced depolarizations in control and CFA neurons. To eliminate the contribution of depolarizations attributable to activation of voltage-dependent Na channels, TTX and a depolarized prepulse are used to block and inactivate Na channels (Ogata and Tatebayashi, 1993; Rush et al., 1998). Under such conditions, we found that, in contrast to ATP-induced subthreshold depolarizations in control neurons, ATP evokes large (>30 mV) depolarizations that exceed the action potential threshold. The suprathreshold depolarization induced by ATP after inflammation is compelling evidence that upregulation of P2X receptors may lead to enhanced firing activity in inflamed neurons.

Different types of sensory neurons have been shown to process distinct pain signals from the periphery to the spinal cord (Willis and Coggeshall, 1991). High percentages of cells are found to respond to ATP in in vitro studies of P2X receptors in normal DRGs (Fig. 2) (Krishtal et al., 1988; Bean, 1990; Burgard et al., 1999; Grubb and Evans, 1999). Inflammation does not change the percentage of total cells responding to ATP (control, 89.4%; CFA, 93.8%), nor does it change the percentage of cells exhibiting fast and slow ATP currents (Fig. 2A). Because of the same pharmacological profiles of ATP responses between control and inflamed neurons (Fig. 3), P2X receptor phenotypes expressed in DRGs are not altered by inflammation. Thus, the homomeric P2X3 receptors are likely to mediate the fast-inactivating ATP currents and heteromeric P2X2/3 receptors are likely to mediate the slow-inactivating ATP currents in inflamed neurons. We and others also show that, in normal rats, the diameters of cells responding to ATP with fast responses are in general smaller than those of cells responding to ATP with slow responses (Fig. 2B) (Tsuda et al., 1999, 2000; Ueno et al., 1999). Behavioral studies suggest that fast-desensitizing ATP responses from small capsaicin-sensitive neurons signal heat and nocifensive behaviors, and slow-desensitizing ATP responses from medium capsaicin-insensitive neurons signal mechanical allodynia (Tsuda et al., 2000). Because the cell size distribution for fast or slow ATP responses remains unchanged after CFA treatment (Fig. 2B), various pain signals are likely to be processed differentially by homomeric P2X3 and heteromeric P2X2/3 receptors in distinct populations of inflamed DRG neurons.

All of our studies were conducted in the somata of DRGs in vitro. The roles of P2X receptors on peripheral and central terminals in nociception are therefore inferred. Although the percentage of cells responding to ATP in intact DRG of control rats is much lower (Stebbing et al., 1998), it seems reasonable to assume that upregulation of P2X receptors in the soma would lead to an increase in P2X receptor expression at both terminals. Activation of P2X receptors at central terminals has been shown to enhance the release of glutamate at synapses in the spinal cord (Li and Perl, 1995; Labrakakis et al., 2000; Nakatsuka and Gu, 2001). The proposed mechanisms underlying the synaptic action of P2X receptor include Ca\(^{2+}\) influx through activated P2X receptors (Robertson et al., 2001) and activation of voltage-dependent Ca\(^{2+}\) channels activated by P2X receptor-evoked action potentials (Cook and McCleskey, 1997; Gu and MacDermott, 1997). Our results suggest that ATP-evoked action potential generation is not likely to occur in control neurons. However, it may underlie the action of P2X receptors in inflamed neurons.
Peripheral P2X receptors at peripheral terminals are known to participate in the transmission of nociceptive and non-nociceptive responses (Cockayne et al., 2000; Hamilton and McMahon, 2000; Hamilton et al., 2000, 2001; Souslova et al., 2000; Tsuda et al., 2000). An increased ATP-evoked depolarization as the result of enhanced P2X receptor expression at peripheral terminals could result in sensitization in sensory afferents. This possibility is consistent with the recent studies of pain behaviors in rats. The concentrations of ATP and αβmεATP used in the behavioral studies in normal rats are $\geq 100 \text{mM}$, a range that is too high to be attained endogenously (Hamilton et al., 1999; Tsuda et al., 2000). After inflammation, ATP concentrations required to elicit pain behaviors are reduced 100-fold (Hamilton et al., 1999). The proposed mechanisms for the increase in the ATP effectiveness in nociceptive signaling include large leakage of ATP from injured cells, sensitization of P2X receptors elicited by enhanced release of neuropeptides or $H^+$ from inflamed tissue, and changes in the second-messenger levels (Hamilton et al., 1999, 2001; Paukert et al., 2001). Although these possibilities cannot be dismissed, our results suggest that upregulation of P2X receptors and enhanced ATP responses are the primary reasons for increased behavioral sensitivity in the inflammatory state. With a twofold to threefold increase in ATP responses after inflammation, a small amount of ATP release would evoke depolarizations large enough to elicit action potentials in DRG neurons (Fig. 7). It is therefore conceivable that endogenous ATP release does not produce pain in normal rats. The same ATP release after inflammation, however, will sensitize neurons and produce abnormal nociceptive responses. Therefore, the profound changes in P2X3 and P2X2/3 receptor expression and in ATP responses observed here may be critical for the induction of pain hypersensitivity after the development of inflammation.

REFERENCES