Responsiveness of C Neurons in Rat Dorsal Root Ganglion to 5-Hydroxytryptamine-Induced Pruritic Stimuli In Vivo

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INTRODUCTION

Itching is an unpleasant sensation that can occasionally degrade the quality of life. There are some similarities between itching and pain; both are unpleasant sensations and most pruritogens, such as histamine, can also produce pain (Dray 1995; Schmelz et al. 2003). The behavioral response in rats or mice to these two types of stimuli, however, has some differences. Itch stimuli evoke scratching or biting, whereas noxious stimuli evoke a withdrawal reflex, flinching, or licking (Kuraishi et al. 1995, 2008; Nojima et al. 2004). Opioids, which are commonly used for alleviation of pain, elicited itching (Hales 1980; Jeon et al. 2005; Maxwell et al. 2005; Slappendel et al. 2000); conversely, the opioid receptor antagonist naloxone enhances pain but inhibits the itch sensation (Metze et al. 1999; Robertson et al. 2008).

To elucidate the characteristics of itch-sensing primary affrents, electrophysiological recordings of these afferents have been made. Microneurography reveals that histamine applied to human skin elicits low-frequency activity in polymodal C fibers (Handwerker et al. 1991; Torebjörk 1974; Van Hees and Gymbel 1972). Tuckett and Wei recorded the activity of A and C fibers in cats and found that only polymodal C fibers were activated by the application of cowhage, which is pruritic in humans (Tuckett and Wei 1987a,b). Polymodal C fiber that responds to cowhage is also reported in monkeys (Johane et al. 2008) and humans (Namer et al. 2008). Histamine-sensitive polymodal C fibers are also responsive to mustard oil, which is known to activate TRPA1, and there is no significant difference in the discharge patterns caused by itching or burning stimuli (Handwerker et al. 1991). A new subtype of C fibers that is insensitive to mechanical stimuli was reported to respond to itch stimuli (Schmelz et al. 1997). However, these fibers also respond to algogens such as capsaicin or bradykinin (Schmelz et al. 2003).

Several chemicals have been used experimentally to elicit the itch sensation. However, the itch-inducing potency of these chemicals differs between species. Histamine, for instance, is a well-known pruritogen in humans. It evokes scratching in ICR mice, hairless guinea pigs, and monkeys (Inagaki et al. 2001; Johane et al. 2008; Woodward et al. 1995), but it is generally less effective in ddY mice and Sprague–Dawley rats (Jinks and Carstens 2002; Kuraishi et al. 1995). Instead of histamine, 5-hydroxytryptamine (5-HT) is often used in mice and rat itch models (Inagaki et al. 2001; Nojima and Carstens 2003a; Nojima et al. 2003; Thomsen et al. 2001; Yamaguchi et al. 1999). Interestingly, 5-HT is only weakly pruritic for humans (Schmelz et al. 2003). 5-HT is released from aggregated platelets and mast cells in rats (Weisshaar et al. 1997). 5-HT produces itching when applied to the human skin (Weisshaar et al. 1997) and is considered to be a cause of pruritus accompanied by polycythemia vera (Fitzsimons et al. 1981). In rats, intradermal injection of 5-HT into the rostral back evokes a scratch response and this behavior is depressed by subcutaneous injection of the opioid receptor antagonist naltrindole (Nojima and Carstens 2003a; Nojima et al. 2003). These observations indicate that 5-HT is a potent pruritogen in rats and suitable for clarifying the properties of afferent fibers conveying the itch sensation to the spinal cord.

In the present study, we intended to answer the following questions. 1) What types of primary afferents are responsible for the reaction to topically applied 5-HT on a receptive field? and 2) What is the characteristic feature of itch-sensing neurons? To address these questions, we made intracellular recordings from rat dorsal root ganglion (DRG) neurons in vivo and...
analyzed the response to itch stimulation induced by topical application of 5-HT to the skin.

**METHODS**

All experimental procedures involving the use of animals were approved by the Committee on the Ethics of Animal Experiments, Kyushu University, and were in accordance with the Guidelines of the Japanese Physiological Society. All efforts were made to minimize animal suffering and the number of animals used for the studies. At the end of the experiments, the rats were given an overdose of urethane and then exsanguinated.

**Behavioral experiments**

In all, 43 male Sprague–Dawley rats (aged 6–9 wk) were used. They were housed under controlled temperature and light. Food and water were freely available. The hair of the rostral back, right hindlimb, thigh, and hip was clipped one day before the behavioral experiments. Before the experiments, the animals were placed in an acrylic cage for about 1 h for acclimation. Behavioral changes were recorded by digital video camera for 1 h. On the clipped rostral back, 5-HT (1% [47 mM] in 99% ethanol, 50 μl) and the vehicle (99% ethanol, 50 μl) were applied on the target skin by using a micropipette instead of being injected intradermally, to avoid eliciting pain by inserting a needle and expanding the skin. The number of scratches to the application site by the hindpaw was counted. We used ethanol as the vehicle. Ethanol can activate TRPV1 and is painful on a skin wound. However, we did not believe it would be painful in this study for two reasons. One is that topical application of ethanol is often used in studies on humans and rarely evokes pain (Ham et al. 2006; Namer et al. 2008; Wasner et al. 2004) and the other is that the amount of penetration of topically applied ethanol to the skin is small (Pennington et al. 2001). We also applied 1% 5-HT (50 μl) to the right hindpaw and observed biting behavior since the electrophysiological experiments were performed from L4 to L6, mainly L5 DRG neurons, which innervate the hindpaw, thigh, and hip. In mouse itch models, biting behavior was elicited by 5-HT injection or under the condition of chronic dermatitis or dry skin (Kuraishi et al. 2008; Maekawa et al. 2002; Nojima et al. 2004). In contrast, a formalin application, which is known to initiate pain behavior, causes licking in mice (Abbott et al. 1995; Hunskaar et al. 1985; Tjølsen et al. 1992). Thus the biting behavior is considered as itch-related behavior. In some experiments, rats were given a subcutaneous injection of naloxone (1 mg·kg⁻¹) or saline at a volume of 1 ml·kg⁻¹ into the back skin 15 min before the topical application of 5-HT.

**Electrophysiological recordings**

In all, 88 rats were used for the electrophysiological experiments. After anesthesia of the rats with urethane (1.2 g·kg⁻¹, administered intraperitoneally), laminectomy was performed at the lumbar level and the right DRG (L4–L6, mainly L5) was carefully exposed with a rongeur. Animals were fixed rigidly in a stereotaxic apparatus and then the skin flaps were stretched by nylon fibers to make a pool for subcutaneous injection of drugs. Subsequently, we gently applied the vehicle (ethanol 20 μl) and then 5-HT (serotonin hydrochloride 20 μl; Sigma) on the receptive field by using a micropipette. We observed the activity of the neurons ≥2 min after applying the drugs to evaluate whether they were responsive to the drugs. Capsaicin (0.05% in ethanol 20 μl; Wako) was applied to the receptive fields of eight 5-HT-insensitive neurons. We did not test the effect of capsaicin on 5-HT-sensitive neurons, since we could not exclude the possibility that previously applied 5-HT would redissolve into the capsaicin solution and activate the 5-HT receptors. C neurons that had membrane potential more positive than −50 mV were excluded from further analysis; however, the firing response of those neurons to 5-HT application or mechanical stimuli was included in the present study due to the difficulty in obtaining recordings from C neurons. Neurons showing spontaneous firing without 5-HT or mechanical stimuli were discarded.

**Statistical analysis**

Total scratching and biting was evaluated by unpaired t-test. Electrophysiological data were evaluated by Mann–Whitney U test or Kruskal–Wallis H test. P < 0.05 was considered significant.

**RESULTS**

**Scratching and biting behavior in response to topically applied 5-HT**

Topically applied 5-HT to the rostral back evoked scratching behavior, whereas the vehicle did not (Fig. 1, A and B; n = 8 in each group). The scratching began within 5 min after application of 5-HT and reached a peak within 10 min; the scratching frequency reached 30 scratches per 5 min and then gradually decreased. The scratching behavior lasted for >40 min. There was a significant delay in the beginning of scratching after 5-HT application, probably due to the diffusion time for 5-HT to reach the nerve terminals. The mean total number of scratch bouts in 1 h induced by 5-HT was 197 ± 27 (mean ± SE, n = 8). Following subcutaneous injection of naloxone (1 mg·kg⁻¹), the scratching was reduced to less than one third (Fig. 1C; vehicle 172 ± 38, n = 8; naloxone 51 ± 18, n = 7), which was statistically significant (P < 0.01), suggesting that the scratching is due to itch but not pain sensation.
Before performing an intracellular recording analysis, behavioral changes induced by the topically applied 5-HT from the sole to the ankle of the right hindpaw were tested. Rats exhibited biting behavior to the 5-HT-applied region but not to the vehicle-applied region (Fig. 1D; n = 6 each group). The biting time course was similar to the scratching time course (Fig. 1E). These observations further indicate that 5-HT applied at the hindpaw, thigh, or hip also initiates itch sensation in rats.

Response of C neurons to topically applied 5-HT

Intracellular recordings were made from 29 neurons with Aβ, 25 neurons with Aδ, and 91 neurons with C-fiber in L5 DRG in vivo (Table 1). Receptive fields of the recorded neurons were identified by applying mechanical or electrical stimulation to the skin. Subsequently, ethanol (99%, 20 μl) or 5-HT (1%, 20 μl) was applied to the receptive field. Neither Aβ nor Aδ neurons responded to topical application of 5-HT (Table 1). Twenty-five of the 91 C neurons (27%) showed orthodromic firing in response to 5-HT application (Table 1). Vehicle application did not produce firings, except one Aδ neuron and one C neuron, which might be cool-sensitive neurons because they responded to the gentle application of ice.

Figure 2 shows an example of a 5-HT-sensitive C neuron (Fig. 2, A–D were recorded from the same neuron). This neuron was also sensitive to touch (arrows) and pinch stimuli (Fig. 2, A and B). Continuous firing began 32 s after topical application of 5-HT (Fig. 2, A and D) and was recorded for >50 min (Fig. 2D). As noticed from the continuous recording (Fig. 2A), action potential was followed by just a small-amplitude afterhyperpolarization (AHP) (Fig. 2C). We classified these neurons as a long-lasting type. These neurons continued to fire for a long time and the peak firing frequency was observed >5 min after 5-HT application. Twelve of the 91 C neurons (13%) were considered to be the long-lasting type and were responsive to noxious stimuli, except one neuron that responded to nonnoxious stimuli (Table 1). We found another type of 5-HT-sensitive C neuron that responded to topically applied 5-HT; however, unlike the long-lasting type, the peak firings were found within 5 min and most of them stopped firing within a few minutes (Fig. 3A). These neurons were sensitive to pinch stimuli (Fig. 3B). We classified these neurons as a transient type and they were also responsive to mechanical stimuli, except for one neuron that responded to nonnoxious stimuli (Table 1). Thirteen C neurons (14%) were considered to be the transient type (Table 1). This type also had a small-amplitude AHP. Figure 4 shows an example trace of a 5-HT-sensitive C neuron. This type of neuron was sensitive to pinch stimuli (Fig. 4B). The 5-HT-sensitive C neurons (73%) were largely sensitive to noxious stimuli, but not to nonnoxious stimuli. A substantial number of 5-HT-sensitive C neurons (11%) were insensitive to mechanical stimuli. 5-HT-sensitive neurons exhibited a relatively large-amplitude AHP, as shown in Fig. 4C.

TABLE 1 Summary of the recorded neurons

<table>
<thead>
<tr>
<th>Factor</th>
<th>5-HT-Sensitive</th>
<th>Aβ neurons (n = 29)</th>
<th>Aδ neurons (n = 25)</th>
<th>C neurons (n = 91)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Transient</td>
<td>Long-Lasting</td>
<td></td>
</tr>
<tr>
<td>Touch</td>
<td>17 (59%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Pinch</td>
<td>11 (38%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Mechanically insensitive</td>
<td>1 (3%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>29 (100%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Touch</td>
<td>3 (12%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Pinch</td>
<td>19 (76%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Mechanically insensitive</td>
<td>3 (12%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Total</td>
<td>25 (100%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Touch</td>
<td>56 (62%)</td>
<td>12 (13%)</td>
<td>11 (12%)</td>
<td></td>
</tr>
<tr>
<td>Pinch</td>
<td>10 (11%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Mechanically insensitive</td>
<td>66 (73%)</td>
<td>13 (14%)</td>
<td>12 (13%)</td>
<td>0</td>
</tr>
</tbody>
</table>
Firing properties of 5-HT-sensitive neurons in response to 5-HT application

Next, we compared the firing properties of long-lasting and transient-type 5-HT-sensitive C neurons. The long-lasting type showed a continuous increase in the number of firings 5 min after the application of 5-HT (Fig. 5A). On the other hand, the number of firings in the transient type reached a peak within 3 min (median 2.5 min, range 1–3 min, n = 13) and then decreased (Fig. 5B). The long-lasting type exhibited a significantly higher firing rate than that of the transient type in the first 5 min (long-lasting type: median 25 spikes/min, range 15–75 spikes/min, n = 12; transient type: median 18 spikes/min, range 3–70 spikes/min, n = 13; P < 0.01; Fig. 5C). Furthermore, the long-lasting-type neurons began to fire much earlier than the transient firing neurons (long-lasting type: median 26 s, range 12–67 s, n = 12; transient type: median 25 s, range 14–115 s, n = 13; P < 0.05; Fig. 5D).

Properties of action potential

The conduction velocity of 5-HT-insensitive and -sensitive C neurons was not significantly different. The median velocity was 0.6 ms⁻¹ in 5-HT-insensitive neurons (range 0.3–1.9 ms⁻¹, n = 66), 0.6 ms⁻¹ in transient-type neurons (range 0.5–1.5 ms⁻¹, n = 13), and 0.5 ms⁻¹ in long-lasting-type neurons (range 0.5–0.7 ms⁻¹, n = 12). There were no significant differences in height or duration of action potential among 5-HT-insensitive, transient, and long-lasting types (height, 5-HT-insensitive: median 83.9 mV, range 55.7–104.5 mV, n = 42; transient: median 80.6 mV, range 67.8–93.9 mV, n = 8; long-lasting: median 75.0 mV, range 56.7–102.7 mV, n = 11; duration, 5-HT-insensitive: median 1.6 ms, range 0.9–3.5 ms, n = 42; transient: median 1.6 ms, range 1.3–2.8 ms, n = 8; long-lasting: median 2.4 ms, range 0.8–4.9 mV, n = 11). Interestingly, the long-lasting type had a significantly smaller fast AHP than that of the 5-HT-insensitive and transient types (Fig. 6, A, B, and C, 5-HT-insensitive: median 7.8 mV, range 0–16.5 mV, n = 42; transient: median 3.0 mV, range 0–10.0 mV, n = 8; long-lasting: median 0 mV, range 0–5.3 mV, n = 11) and they were significantly different. The long-lasting-type neurons also showed no apparent slow AHP following the fast AHP, even though the action potential had a large Ca²⁺ component in the falling phase.

Application of algogenic agents to 5-HT-insensitive C neurons

To clarify whether 5-HT-insensitive C neurons were responsive to algogenic stimuli, we topically applied capsaicin...
(0.05% in ethanol) to the receptive fields (Fig. 7, A and B). Before capsaicin application, 5-HT was topically applied to test whether a neuron responded to 5-HT (Fig. 7A) and then noxious pinch stimulation was added. The neuron shown in Fig. 7A responded to pinch stimuli but not to 5-HT. In the same neuron, topical application of capsaicin elicited continuous firing lasting for >5 min with long latency (median 62 s, range 46–77 s, n = 2). Two of eight neurons were responsive to capsaicin.

Location of the receptive fields

Figure 8 shows the location of the receptive fields that we recorded. They were located mostly on the hindpaw and hip. The distribution of long-lasting C neurons, transient C neurons, and 5-HT-insensitive C neurons did not vary.

DISCUSSION

In the present study, intracellular recordings were made from C neurons in rat DRG in vivo to elucidate the firing properties of neurons in response to topically applied 5-HT to the skin, which elicited an itch-associated scratching response in rats. A small population of C neurons responded to the topical application of 5-HT and these 5-HT-responsive C neurons could be divided into two subtypes based on their firing duration, firing rate, and latency of the response. One type of C neurons showed long-lasting firing, higher firing frequency, and short latency. The firing duration was comparable to the behavioral changes induced by 5-HT topically applied to the hindlimb. The other type of C neurons exhibited transient firing that lasted for <5 min, low firing frequency, and long latency. These observations suggest that the long-lasting-type C neurons may express 5-HT receptors on the peripheral nerve endings and play an important role in carrying the itch sensation to the spinal cord in rats. The other possibility is that the shorter delay and longer duration of the 5-HT response could also be due to a higher density of 5-HT receptors.

Several chemicals have been reported to induce the itch sensation, including histamine, 5-HT (Nonjima and Carstens 2003a,b; Thomsen et al, 2001; Yamaguchi et al. 1999), substance P (Andoh et al. 1998), trypsin (Uj et al. 2006), and prostaglandin E2 (Neisius et al. 2002). Histamine is the best-known itch mediator in humans, but not in rodents (Kuraishi et al. 1995), whereas 5-HT is a potent pruritogen in rats (Thomsen et al. 2001). In the majority of behavioral studies, 5-HT is administered by intradermal injection, whereas in the present study, topical application was used, which has the advantage of avoiding mechanical pain stimulation induced by intradermal injection. Similar to the results obtained in previous studies, the topical application of 5-HT evoked scratch behavior, which was reduced by prior application of naloxone, known to enhance the sensation of pain. In agreement with this finding, it is well known that intrathecal injection of an opioid initiates an itch sensation that can be blocked by naloxone (Hales 1980; Jeon et al. 2005; Maxwell et al. 2005; Slappendel et al. 2000). When 5-HT was injected into the hindpaw, rats bit the injected region instead of scratching, since it is difficult for them to scratch their hindpaw (Kuraishi et al. 2008). In the itch model that we used in the present study with topical application of 5-HT to the hindpaw, biting behavior was also observed. In contrast, an algogenic agent, such as formalin, applied to the

The 5-HT-sensitive long-lasting-firing C neurons started firing earlier than the C neurons of the transient type. This suggests that the long-lasting C-afferent terminals exist at a more superficial layer of the skin than those of the transient type. Numerous fine nerve endings are spread out at a level just below the epidermis (Shelly and Arthur 1957) and free nerve endings in the dermoepidermal junction are regarded as a receptor for itch sensation (Wahlgren 1992). It is reported that patients with atopic dermatitis have an increased number of nerve fibers in the skin (Tobin et al. 1992; Urashima and Mihara 1998) and nerve growth factor–mediated sprouting of nerve fiber is found in patients with contact dermatitis (Kinke-lin et al. 2000). These reports support our findings. Interestingly, the onset of scratching response was later than the onset of action potentials of long-lasting neurons. There may be two reasons why scratching appears later: one is that an itchy sensation that produces a behavioral change needs the accumulation of activity of itch-related neurons; the other is that transient-type neurons may transmit painful sensation and they may suppress the itch sensation transiently in the first several minutes. However, we do not have enough evidence at present to make such hypotheses.

In the present study, topically applied 5-HT to the receptive field evoked repetitive firing in a small population of C neurons but not in Aβ and Aδ neurons. Previous electrophysiological observations support the contribution of C-afferent fibers to the itch sensation; for instance, cowhage selectively activates polymodal C fibers in rats and monkeys (Johanek et al. 2008; Tuckett and Wei 1987b) and histamine activates a subset of C fibers in humans (Handwerker et al. 1991; Schmelz et al. 1997, 2003). Recently, it was reported that the histamine-induced itch sensation is mediated by the activation of TRPV1 receptors expressed at C afferents via an arachidonic acid metabolite produced by histamine receptor activation (Shim et al. 2007). These observations are consistent with our findings. However, it is possible that multiple pathways might be responsible for the itch sensation and it is still not clear whether 5-HT-sensitive C neurons belong to the previously reported itch-responsible primary afferents.

From our behavioral study, naloxone inhibited itch-associated scratching behavior. The site of opioid action on the itch sensation pathway has not been identified. It is well known that the μ-receptor agonist DAMGO ([D-Ala, N-Me-Phe, Gly-ol]-

FIG. 6. Comparison of amplitude of AHP among C neurons. A: representative action potential of 5-HT-sensitive long-lasting-type neuron. B: representative action potential of 5-HT-insensitive C neuron. Note that obvious AHP was observed in 5-HT-insensitive C neuron. C: amplitudes of AHP of 5-HT-insensitive neurons (n = 42), 5-HT-sensitive long-lasting neurons (n = 11), and 5-HT-sensitive transient neurons (n = 8) are plotted. Filled circles represent mechanosensitive C neurons and open circles represent mechanoinensitive C neurons. **P < 0.01 (Kruskal–Wallis H test).

FIG. 7. Representative trace of 5-HT-insensitive C neuron that responded to pinch stimuli and topically applied capsaicin. A: 5-HT did not evoke firing. B: continuous firing was elicited following topical application of capsaicin to the same neuron.
enkephalin) applied to the spinal cord inhibits the release of glutamate from the primary afferents, including C fibers (Ikoma et al. 2007; Kohno et al. 1999) and causes membrane hyperpolarization in substantia gelatinosa neurons (Yoshimura and North 1983), indicating that the analgesic effect of opioid is exerted both on the primary afferents and spinal interneurons. In contrast, subcutaneous injection of naltrexone does not affect the activity of cutaneous nerves innervating the chronic dermatitis area (Maekawa et al. 2002). Moreover, neither systemic morphine nor naltrexone affected the 5-HT-evoked c-Fos-like immunoreactivity in the superficial laminae of the spinal dorsal horn, whereas morphine significantly attenuated the intradermal capsaicin-induced immunoreactivity (Nojima et al. 2003). Therefore the μ-opioid receptor may modulate the itch sensation at the supraspinal level; however, further experiments are necessary to reveal how opioid receptors act on itch transmission.

Interestingly, the 5-HT-sensitive long-lasting-firing neurons had a significantly small, fast AHP. In general, nociceptive DRG neurons have a broader action potential than that of low-threshold mechanoreceptive neurons in the same conduction velocity group (Djouhri et al. 1998; Fang et al. 2005; Ritter and Mendell 1992). Nociceptive C neurons have a broader action potential and larger AHP than those of C neurons responsive to nonnoxious stimuli (Djouhri et al. 1998; Fang et al. 2005). The amplitude of AHP observed in this study was smaller than that previously reported (Fang et al. 2005), probably due to the more negative resting membrane potential of C neurons in our study. However, the resting membrane potential in three types of neurons was not significantly different in the present study; therefore the 5-HT-sensitive long-lasting C neurons may have smaller AHP compared with that of 5-HT-insensitive C neurons.

Regarding the issue of whether itch-sensing primary afferents respond to mechanical stimuli, Fang et al. (2005) reported that 54% of C fibers were nociceptors, 11% were nonnociceptors, and 34% were unresponsive to mechanical stimuli. The mechanical threshold of C fiber ranges from 16 to 608 mN (Pogatzki et al. 2002). In the present study, almost all 5-HT-sensitive C neurons responded to mechanical noxious stimuli and several were responsive to nonnoxious stimuli. Thus the 5-HT-sensitive neurons also responded to mechanical stimulation, indicating that the 5-HT-sensitive C neurons are polymodal. In this study, we did not apply thermal stimulation or capsaicin to the receptive fields of 5-HT-sensitive neurons. 5-HT receptor mRNAs were expressed in rat dorsal root ganglion neurons (Nicholson et al. 2003). 5-HT₂A induces thermal hyperalgesia in acute injury and inflammation in rats (Tokunaga et al. 1998). Activation of metabotropic 5-HT receptors enhances the TRPV1 function in primary afferent neurons (Ohta et al. 2006). This evidence suggests that 5-HT-sensitive C neurons might be part of the polymodal neurons. Furthermore, we applied capsaicin to the 5-HT-insensitive neurons, 85% of which were activated by noxious mechanical stimuli, and some of them responded to capsaicin (Fig. 7), indicating that the 5-HT-insensitive primary C neurons include polymodal neurons. It is not known from our study whether 5-HT-responsive C neurons are the only pruritogen-sensitive primary afferents. Endopeptidases elicit a pure itch sensation without any wheal or flare (Arthur and Shelley 1955). It was found that itching could be induced by electrical stimulation to the wrist without generating an axon reflex (Ikoma et al. 2005). It has been reported that histamine and another typical itch agent, cowhage, activate separate populations of spinothalamic tract neurons (Davidson et al. 2007), implying that the itch sensation is not mediated by a single system. Recently, it was reported that gastrin-releasing peptide receptors mediate itch sensation in the spinal cord (Sun and Chen 2007). Gastrin-releasing peptide is expressed in a subset of small and mediumsized DRG neurons that are peptidergic unmyelinated fibers (Sun and Chen 2007). Since it is still unknown whether long-lasting-type 5-HT-sensitive C neurons express gastrin-releasing peptide or peptidergic markers such as substance P and calcitonin gene-related peptide, further experiments are needed to answer the question.

The 5-HT receptor subtype responsible for activation of the primary afferent is still unknown. In DRG, the mRNA for 5-HT₁B, 5-HT₂A, 5-HT₂B, 5-HT₃B, and 5-HT₄ receptors was detected in small-diameter neurons (Nicholson et al. 2003). It is known that 5-HT₂A and 5-HT₄ receptors have a role in pain sensation (Eschalier et al. 1989; Giordano and Rogers 1989; Okamoto et al. 2002; Tokunaga et al. 1998). In vitro electrophysiological experiments showed that 5-HT modulates the activity of primary afferents. Activation of axonal 5-HT₃ receptors enhances membrane excitability and modulates action potential trains in unmyelinated nerve fibers (Lang et al. 2006). Ohta et al. (2006) showed that 5-HT changed TRPV1 functions through the activation of 5-HT₂A and 5-HT₇ receptors in cultured DRG neurons. It is reported that 5-HT-evoked itching is mediated by 5-HT₇ receptors (Nojima and Carstens 2003b; Yamaguchi et al. 1999); however, this study does not indicate the principal receptor for 5-HT-induced itching. Because the latency of firing after topical application of 5-HT was within 1 min in the majority of 5-HT-sensitive long-lasting C neurons, these data may suggest that 5-HT directly activates 5-HT receptors; however, we cannot exclude the possibility that the activation of long-lasting C neurons was mediated by surrounding tissues such as keratinocytes or mast cells with the release of certain chemicals.

In conclusion, the itch-sensing primary afferents might be polymodal C fibers that responded to topically applied 5-HT, resulting in continuous firing for ≈50 min. This long-lasting high-frequency firing was, in part, due to the small amplitude of fast AHP and the absence of slow AHP, even though the neurons had large Ca²⁺ components in the action potential.
The functional significance of the transient-firing neurons evoked by 5-HT has not been clarified in the present study. In view of the fact that 5-HT is also an algesic substance and considering the long duration of the scratching behavior demonstrated in the present study, we assumed that the transient firing observed in a subpopulation of C neurons may be responsible for carrying pain sensation to the spinal dorsal horn.

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**Disclosures**

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