Tactile C fibers and their contributions to pleasant sensations and to tactile allodynia

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In humans converging evidence indicates that affective aspects of touch are signaled by low threshold mechanoreceptive C tactile (CT) afferents. Analyses of electrophysiological recordings, psychophysical studies in denervated subjects, and functional brain imaging, all indicate that CT primary afferents contribute to pleasant touch and provide an important sensory underpinning of social behavior. Considering both these pleasant and social aspects of gentle skin-to-skin contact, we have put forward a framework within which to consider CT afferent coding properties and pathways—the CT affective touch hypothesis. Recent evidence from studies in mice suggests that CTs, when activated, may have analgesic or anxiolytic effects. However, in neuropathic pain conditions, light touch can elicit unpleasant sensations, so called tactile allodynia. In humans, tactile allodynia is associated with reduced CT mediated hedonic touch processing suggesting loss of the normally analgesic effect of CT signaling. We thus propose that the contribution of CT afferents to tactile allodynia is mainly through a loss of their normally pain inhibiting role.

Keywords: touch, unmyelinated, tactile allodynia, fMRI, psychophysics, social

Historically, human tactile sensibility was considered to be mediated solely by low-threshold mechanoreceptors with large myelinated (Aβ) afferents conducting impulses at high speed (around 50 m s⁻¹). In contrast, unmyelinated low-threshold mechanoreceptive afferents (C-LTMRs) have been known to exist in the hairy skin of mammals since 1939 (Zotterman, 1939; Douglas and Ritchie, 1957; Bessou et al., 1971; Iggo and Kornhuber, 1977; Kumazawa and Perl, 1977). For long, it was assumed that humans did not share this seemingly primitive tactile system with other mammals. Nevertheless, in recent years it has been demonstrated repeatedly that human skin is also innervated by C-LTMRs conducting impulses with a speed of only about 1 ms⁻¹. In man, these nerve fibers were first found in microneurography recordings from the infra- and supra-orbital nerves (Johansson et al., 1988; Nordin, 1990). Soon after, they were found in the arm and leg suggesting a more general distribution (Valbo et al., 1993, 1999; Edin, 2001; Wessberg et al., 2003; Campero et al., 2011). In humans, C-LTMRs are called C tactile or CT afferents but so far afferent response properties seem to be similar across species (Valbo et al., 1999).

Although there is currently no accurate method to assess the innervation density of CT afferents in humans, it is a recurring experience in microneurography recordings from the lateral ante-brachial cutaneous nerve of the forearm that they are encountered as often as Aβ afferents. CT afferents have never been found in the palm of the hand despite numerous microneurography recordings from this skin area.

C TACTILE (CT) AFFERENTS

CT afferents respond to indentation forces in the range 0.3–2.5 mN (Valbo et al., 1999), tested with von Frey monofilaments, and are thus as sensitive to skin deformation as many of the Aβ afferents. CT afferents respond with high frequency to stimuli that are clearly innocuous, such as slow stroking with the experimenter’s finger tips or a soft brush (Figures 1A–C; Valbo et al., 1999). In contrast to C nociceptors, with mechanical thresholds >2.5 mN, CT afferents do not distinguish between pin pricks and smooth probe indentations but respond equally well to both these types of stimuli. C nociceptors may also respond to light brush stroking but their responses never exceed a few impulses (Valbo et al., 1999).

The conduction velocity of CT afferents, as assessed with mechanical or electrical stimulation, varies between 0.6–1.3 ms⁻¹. To a sustained indentation, CTs initially respond with a high frequency burst of impulses but the firing rate decreases to zero within 5 s. The adaptation characteristic of CT afferents is thus intermediate in comparison with the slowly and rapidly adapting myelinated mechanoreceptors; slowly adapting units continue to fire during indentation whereas rapidly adapting units only fire when the skin deformation is changing. In a subset of CT afferents the response may increase again after the initial period of adaptation with firing continuing for 1–2 min until it finally stops; a phenomenon described as delayed acceleration (Valbo et al., 1999). A related phenomenon has been described for rat nociceptors (Andrew and Greenspan, 1999). Another feature of CT afferents is that they are highly fatigable. When several identical
stimuli are delivered to the same skin area the response to the first stimulus is usually much larger than the following responses to identical stimuli. When a skin deformation is released CT afferents may produce after-discharges that may last up to several seconds (Nordin, 1990).

The receptive field of a human CT afferent is roughly round or oval in shape with no preferred orientation. Detailed analyses has revealed that, in humans, the field consists of 1–9 small responsive hot spots distributed over an area up to 35 mm$^2$ (Figure 2; Wessberg et al., 2003).

A caressing type of slowly moving touch is a particularly effective stimulus for CT afferents. It has been measured through single unit microneurography that the maximal unit response occurs for movement velocities in the range 1–10 cm s$^{-1}$ whereas the response is weaker for slower and faster movements (Loken et al., 2009; Figure 3A). In psychophysical experiments brush stroking in the same velocity range 1–10 cm s$^{-1}$ is perceived as more pleasant than stroking with slower or faster velocities (Figure 3B). Indeed, there is a positive correlation between firing frequency of CT afferents and perceived pleasantness of soft brush stroking (Figure 3C).

**FINDINGS IN SUBJECTS LACKING LARGE MYELINATED (Aβ) AFFERENTS**

Direct evidence for a specific role of CT afferents in tactile sensation has been difficult to acquire; a major reason being that it is not possible to stimulate CT afferents without also activating Aβ afferents. Unique data has been collected from two subjects selectively lacking Aβ afferents but who have intact C fibers as the result of sensory neuronopathy (a rare disorder of nerve cell bodies of the large primary sensory neurons) (Sterman et al., 1980). The two subjects (initials GL and IW) are well described in the literature (Forget and Lamarre, 1987; Cole and Sedgwick, 1992). They have been studied extensively over the years particularly with regard to motor functions because of their proprioceptive deficit. It had also been reported, although merely in passing, that they had lost all tactile sensations when they became ill. This observation was consistent with the view at that time that tactile sensation was altogether dependent on Aβ signaling was largely based on nerve block experiments in healthy subjects demonstrating a lack of tactile sensations when Aβ fibers were blocked through pressure applied on the nerve (Mackenzie et al., 1975).

When it became evident that human skin is supplied with a system of unmyelinated afferents, it became necessary to re-examine the tactile sensibility of these rare neuronopathy subjects using more refined approaches. Rigorous psychophysical tests were pursued to explore if the neuronopathy subjects were able to detect CT targeted touch. It was found that subjects lacking Aβ afferents detected soft brush stroking and weak monofilament indentation on the forearm skin where CT afferents are abundant (Olausson et al., 2002, 2008; Cole et al., 2006). Importantly, they failed altogether to detect the same kind of stimuli applied to the glabrous skin of the hand where CT afferents are lacking. In addition, they were unable to detect vibratory stimuli which give a poor activation of CT afferents but a vigorous activation of Aβ afferents (Iggo, 1960; Bessou et al., 1971; Kumazawa and Perl, 1977; Olausson et al., 2002, 2008).
The sensation reported by the patients in association with massive and selective CT input (soft brush stroking of the hairy skin) was weak, vague, and inconsistent. In some trials the subject reported no sensations at all. In others, they reported a sensation of light touch which was barely detectable and difficult to describe. One of the subjects (GL) reported that she began to feel more touch sensations in her daily life once she had had the experience of touch perception from the affected skin areas during the experiments and had become aware of this type of perceptual experience. Although the two neuronopathy subjects were not able to give a concise or detailed description of the sensation elicited by CT stimulation, they both reported, independent of each other, that it was a pleasant touch experience with no hint of pain, tickle, or itch. None of the two neuronopathy subjects feel tickle in the affected skin areas which contradicts the old hypothesis that CTs may signal a tickling sensation (Zotterman, 1939; but see Fukuoka et al., 2013). The neuronopathy subjects’ ability to spatially localize CT stimulation is very poor; they make mistakes when trying to identify which body quadrant is being stimulated, although they overall perform above chance level (Olausson et al., 2008).

**Findings in Subjects Lacking C Afferents**

We have also examined patients with a hereditary disorder associated with a nerve growth factor beta (NGFB) gene mutation causing a denervation pattern opposite to that of the neuronopathy subjects GL and IW. Carriers of the NGFB mutation show a reduction in density of thinly myelinated and unmyelinated nerve fibers, thus likely including CT afferents, whereas their Aβ afferents are intact. Their condition has been classified as hereditary sensory and autonomic neuropathy type V (HSAN-V). We have addressed the relationship between C fiber function and pleasant touch perception in 10 HSAN-V individuals from a unique population of carriers (Morrison et al., 2011). The HSAN-V patients perceive gentle, slow stroking, optimal for eliciting CT afferent responses (1–10 cm s⁻¹), as less pleasant than do matched controls and also differ in their rating patterns across stimulation velocities. Hence, these observations further support the notion that CT afferents make a critical contribution to the perception of affective touch.

**Cortical Processing of C Tactile (CT) Stimulation**

When functional magnetic resonance imaging (fMRI) is used to study brain responses to touch stimuli in neurologically intact subjects and in neuronopathy subjects lacking Aβ afferents, different sensory areas are activated by Aβ and CT afferents. In healthy subjects soft brush stroking activates the classical somatosensory areas S1 and S2 as well as insular cortex, notably the posterior part of the contralateral insular cortex (Olausson et al., 2002). When similar brushing stimuli are applied to the neuronopathy subjects lacking Aβ afferents (GL and IW) only the posterior insular region is activated (Olausson et al., 2002, 2008; Figure 4). Further, there is a somatotopic organization of CT responses in the posterior insular cortex with forearm projecting anterior to thigh stimulation (Björnsdotter et al., 2009). The somatotopic arrangement suggests that CT afferents follow the thin-fiber spinothalamic pathway with the posterior insular cortex as the primary cortical receiving area (Craig, 2002). No corresponding insular activation was found for brush stroking in the C-fiber denervated HSAN-V subjects (Morrison et al., 2011).

CTs have not been found in the glabrous skin of the hand, yet it is commonly observed that glabrous skin touch is also perceived as pleasant. When contrasting the brain activation of slow brush stroking on the forearm to that of slow brush stroking in the palm there is a significantly greater activation of the posterior insular cortex and mid-anterior orbitofrontal cortex (OFC) for brush stroking on the hairy skin of the forearm (McGlone et al., 2012). The opposite contrast (stroking on the arm minus stroking in the palm) shows a significant activation of somatosensory cortices. Although psychophysical ratings show no differences in intensity or pleasantness ratings, a touch-questionnaire in which subjects used a newly developed “Touch Perception Task”.

![FIGURE 3](https://example.com/figure3.png)
that is likely to signal close, affiliative body contact with others (Morrison et al., 2010).

**C TACTILE (CT) AFFERENTS AND TACTILE ALLODYNA**

Tactile alldynia is a symptom of neuropathic pain where normally innocuous moving tactile stimuli produce pain. People with tactile alldynia typically experience a burning, tender sensation during soft stroking of the affected skin (Rasmussen et al., 2004). Even a very light stimulus, such as a patient’s garment brushing against the skin during movement, can evoke alldynia. The prevailing hypothesis for tactile alldynia is changed tactile signaling in the spinal cord (Woolf, 1993) following central sensitization where Aβ LTMRs signal to nociceptive neurons in the dorsal horn and from there to cerebral pain processing areas (Campbell et al., 1988; Torebjork et al., 1992; Woolf, 1993; Wasner et al., 1999). This view is based on human selective nerve block experiments demonstrating that tactile alldynia is abolished by compression or ischemic block of Aβ afferents (Gracey et al., 1992; Torebjork et al., 1992; for contradictory results see Nagi et al., 2011).

The view of a critical role for Aβ afferents in mediating human tactile alldynia was established at a time when C-LTMRs were generally thought not to exist in humans. The first study to suggest a critical role for C-LTMRs in signaling alldynia used a vesicular glutamate transporter type 3 (VGLUT3) knock-out mouse, which functionally disconnects signaling from C-LTMRs by preventing glutamate release (Seal et al., 2009). After the loss of VGLUT3 neurons mechanical hypersensitivity following inflammation, nerve injury and trauma is reduced, thus suggesting a critical role for C-LTMRs in mechanical hypersensitivity (Seal et al., 2009). Furthermore, electrophysiological recordings in rats demonstrate a possible anatomical pathway for tactile alldynia where C-LTMRs project to lamina I spinoparabrachial wide dynamic range neurons (Andrew, 2010). Nevertheless, a later study found that preventing the development of C-LTMRs resulted in mice with no hypersensitivity (Lou et al., 2013).

Recently, new light has been shone on this question through the identification of the C-LTMR specific marker TAF4 (Delfini et al., 2013). Following inflammation and nerve injury TAF4 knock-out mice show enhanced mechanical and chemical hypersensitivity, and this effect is reversed by application of the TAF4 protein (Delfini et al., 2013). The authors speculate that upon activation, C-LTMRs might release both glutamate and TAFA4 with glutamate promoting mechanical hypersensitivity and TAFA4 instead preventing mechanical hypersensitivity. This suggestion also provides a potential explanation for the different findings regarding the functional knock-out of glutamate signaling (Seal et al., 2009) and the complete loss of C-LTMRs (Lou et al., 2013). Losing glutamate alone as in the study by Seal et al. would leave TAFA4 unopposed and drive resistance to hypersensitivity. However, in the case of a complete loss of C-LTMRs (Lou et al., 2013) both glutamate and TAFA4 would be lost leaving no net change in sensitivity. But there are other potential explanations for the discrepancy between these two studies. For example, in Lou et al. experiments there is a disrupted development, so the results might reflect compensation for growing up without C-LTMRs. Alternatively, spinal cord and brain neurons, which also express VGLUT3, may mediate the injury-induced
hypersensitivity seen in Seal et al. experiments, rather than C-LTMRs.

A pain modulatory role for C-LTMRs was suggested earlier in a study in rats indicating that C-LTMR targeted input may inhibit C-nociceptive messages in the dorsal horn (Lu and Perl, 2003). By conducting electrophysiological experiments, a specific inhibitory pathway was identified between substantia gelatinosa neurons receiving direct peripheral C-LTMR afferent projections and other substantia gelatinosa cells receiving direct nociceptive input (Lu and Perl, 2003). This unmyelinated circuit represents a potential pathway for C-LTMR impulses to suppress nociceptive impulses (Lu and Perl, 2003). Further, in wild-type mice administration of TAF4a reverses the effect of injecting an inflammatory agent (carrageenan) normally causing mechanical hypersensitivity, consistent with an analgesic role for C-LTMRs (Delfini et al., 2013).

The topic of C-LTMRs in pain inhibition also ties back to the finding of pharmacogenetic activation of MRGPRB4+ expressing neurons (thought to be C-LTMRs) promoting conditioned place preference in mice, indicating that such activation is positively reinforcing and/or anxiolytic (Vrontou et al., 2013), mechanisms which also may have a role in pain modulation.

Based on this animal literature we set out to examine the contribution of CT afferents to the allodynic condition in humans using the heat capsaicin model of dynamic tactile allodynia (Liljencrantz et al., 2013). The contribution of CT afferent signaling was addressed by studying healthy subjects as well as the two rare patients with selective denervation of Aβ afferents (GL and IW). Following application of the model healthy subjects reported tactile evoked pain whereas the patients did not. Instead, both subjects spontaneously reported that the stroking sensation from the allodynic zone was different to their C-touch sensation (faint sensation of pleasant touch) familiar to both subjects. When asked to further describe how the sensation differed, they both, independent of each other, said “weaker sensation” for stimuli from the allodynic zone compared to a control area. Since the OFC is a key area for CT hedonic processing (McGlone et al., 2012; cf. above) these findings suggest that dynamic tactile allodynia is associated with reduced CT mediated hedonic touch processing. Nevertheless, since the patients do not develop allodynic pain (Treede and Cole, 1993; Liljencrantz et al., 2013), this seems dependent on Aβ signaling, at least under these experimental conditions.

Considering a possible analgesic effect of C-LTMR signaling (Lu and Perl, 2003; Delfini et al., 2013; Vrontou et al., 2013) it seems pertinent to speculate that in neuropathic pain conditions there is a gating resulting in a loss of the pain inhibition mediated by C-LTMRs to prioritize nociceptive signaling. This would be consistent with the canonical view that tactile allodynia is signaled by Aβ afferents, and we thus propose that the contribution of C-LTMR/CT afferents is mainly through a loss of their normally pain inhibiting role.

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