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Azanon Gracia, Elena and Longo, Matthew R. (2019) Tactile perception: beyond the somatotopy of the somatosensory cortex. *Current Biology* 29 (9), R322-R324. ISSN 0960-9822.

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## **Tactile perception: Beyond the somatotopy of the somatosensory cortex.**

Elena Azañón & Matthew R. Longo

### **Short abstract:**

New research demonstrates systematic errors of tactile localisation, involving confusions of body parts and body sides. Such errors do not follow the organisation of topographic maps in somatosensory cortex, suggesting that tactile localisation involves coding of abstract features of limbs.

### **Text:**

The skin can convey many sensations, which are translated into our ability to perceive, for instance, the smooth texture of a marble statue or the pinprick of a mosquito bite. Common to all these sensations is the phenomenological experience that they occur somewhere on the body. Indeed, unlike other bodily sensations such hunger or fatigue, it is difficult to conceive of a tactile sensation which can be felt but not localized, whether in its current true spatial location or in an erroneous one [1]. Even in the extreme case of patients who are unable to locate tactile sensations correctly, touches are not perceived floating in the air, but as happening somewhere within the confines of their body [2,3]. Touches on the body, are initially represented in the cortex in an orderly manner, so that adjacent parts of the skin are represented by adjacent bits of the brain [4]. This has been famously illustrated in textbook depictions of Penfield's sensory homunculus in the primary somatosensory cortex (S1; Figure 1). Given this cortical topography, one might assume that the problem of localising a touch on the skin reduces to localising a corresponding peak of activation in these somatotopic

maps. Indeed, several studies have shown that the brain can represent tactile location via population coding within somatotopic maps [5], and some reports of mislocalizations in amputees suggest a dependency on homuncular organization, as when touches on the ipsilateral face, which is adjacent to the arm in the homunculus, evoke tactile sensations on the phantom arm [6]. A new study by Badde et al. [7] in this issue of *Current Biology*, challenges this “homuncular” view and delivers important new insight into how our nervous system relies on abstract features or categorical information to locate touch.

Existing neurocognitive models of somatosensory processing have suggested that localizing a touch within a somatotopic map is not by itself sufficient to localize the stimulus on the body surface, but require higher order body representations (e.g., [8]). In the early 20<sup>th</sup> century, before the discovery of the topographical architecture of the somatosensory cortex [4], Head and Holmes [3], already proposed a mental representation of the body surface to account for the deficits of patients who could detect touch, but were unable to localize where on the body the touch was. This representation has come to be called the superficial schema. We do not have to go so far from our daily routines, however, to envision the need for such representation. Now spare a thought for the kid who is learning to play the guitar. With training, the cortical representation of the fingertips of her left hand which she uses for fingering the strings on the fretboard will enlarge as compared to other kids of her age (cf. [9]). But, how after such cortical changes is she still able to localize touch without producing massive mislocalizations? This is where a body representation, such as the superficial schema comes into play, as a link between cortical maps and body locations that is updated to reflect plastic changes to those maps, and that accounts for their inherent distortions and magnifications [8]. Nevertheless, such models have difficulty accounting for puzzling mislocalizations observed in some amputees, where tactile stimuli on a foot are occasionally

attributed to the absent hand [6], since the hand and foot are far apart in the somatosensory homunculus (Figure 1).

More than a century after Head and Homes' first insights into tactile localization, Badde et al.'s [7] research provides remarkable evidence for mislocalizations in healthy individuals that are, as in the example with the amputees, not determined either by the sensory homunculus or the superficial schema. They show that when touched on a limb, healthy adults occasionally – but systematically – misattribute the touch to a limb that was not touched at all. Many of these “phantom errors” were attributed to the homologous contralateral limb, e.g., the right hand for a touch on the left hand, a pattern that could possibly be explained by reliance on the somatosensory homunculus given growing evidence of the existence of bilateral receptive fields [10] and inter-hemispheric interactions [11], as early as in S1 (for a review see [12]). However, a smaller set of phantom errors were systematically attributed to opposite limb types, e.g., a foot for a hand and vice-versa, which excludes overlapping representations in S1 as a source of these phantom errors, since the hand and foot are far apart in S1. One possibility could be that these mislocalizations might arise from overlapping representations in S2 (secondary somatosensory cortex), where the foot and hand representations are adjacent, at least in monkeys [13]. However, an explanation based on adjacent representations in S2, could only partially account for the pattern of mislocalizations observed in Badde et al.'s study [7]: while it could explain why phantom errors are more frequent across feet and hands sharing the same body side, it cannot account for the fact that mislocalizations are more often than chance referred to the limb placed at the stimulated limb's canonical side of space (e.g., which for the right hand would be the right side of space).

The mislocalizations reported here are therefore consistent with their resulting from representations in which touch is coded in terms of a number of abstract features, rather than

as a spatial map. This way of representing touch may be related to some other results in the literature. For example, interleaving the fingers of a hand can lead to confusions about what hand is touched, but not which finger, despite the fact that finger type and laterality could both rely on the same activity in S1 [14]. Similarly, the pattern of mislocalizations at the hairy and glabrous skin are virtually identical, despite being clearly different at the level of S1, which suggest the use of a representation abstracting away from the details of each individual skin surface [15]. The pattern of mislocalizations reported by Badde et al. [7] provides new and exciting insight into the type of information that is needed to encode location, and appears to implicate the use of representations based on body-related categorical information. This idea fits with recent studies showing that motor representations of the body in the posterior parietal cortex show substantial degrees of mixed selectivity, with common responses to movements made with the hand and foot, or by the left and right hands (e.g., [16]). The present results suggest that similar mixed coding may also be employed in touch (Figure 1).

A representation based on high-dimensional mixed-selectivity would produce very different patterns of confusions and errors than one based on topographic maps. The phantom errors reported by Badde et al. [7] emphasise three relevant features, in decreasing order of importance: body part (i.e., a hand or a foot), body laterality (i.e., right or left), and canonical spatial location. These illusory percepts resemble those obtained in vision, the so-called illusory conjunctions, in which people report single objects having a combination of features which were not actually presented [17]. Indeed, Badde et al.'s [7] results can be thought of as a form of illusory conjunction on touch based on the features of limbs. Studies of illusory conjunctions in vision show that such conjunctions are modulated by stimulus similarity [17]. The present results suggest that a similar process might take place in touch, and demonstrate at least three relevant categorical features along which similarity is defined. For instance,

phantom errors typically matched the correct limb with respect to limb type, and body side. Interestingly, a third feature is the body part's *canonical* location. Several accounts have highlighted, based on indirect evidence, the role of the canonical location of the limb representation in tactile localization at initial stages of sensory processing. For instance, by assuming that immediately following stimulation, touch is referred to a stored representation of the default posture [8] (for a review see [12]). Badde et al. [7] results provides direct evidence of the existence of such canonical representations.

Perhaps, one of the most surprising points in Badde et al. [7], is the finding that the *external* location of the touched limb did not predict the pattern of mislocalizations across hand and foot. This contradicts the widespread idea that touch takes the posture of the touched limb into account even in situations where there is no overt reaction to touch (e.g., [18]). On the contrary, Badde et al. [7], propose that when explicit continuous localization is not needed, for instance to report or move the touched limb, tactile processing can abstract away from topographic representations (somatotopic or external), and use bodily related categories relevant to the task itself. This raises a potentially interesting analogy to studies in vision which have argued for two distinct systems for spatial representation, one based on precise spatial coordinates, the other based on categorical spatial relations [19]. A question that remains to be answered, however, is under which circumstances touch relies on categorical vs coordinate spatial relations. For example, the so-called 'crossed-hands deficit', in which the ability to tell which of the hands was touched first is impaired when the hands are crossed, has been widely-used as an experimental model of tactile localisation in external space [20]. The present results, however, raise the possibility that the deficit may not result from the precise coding of touch in continuous external space, but from an incorrect assignment of touch to a particular limb.

The results of Badde et al. raise fundamental questions about our experience of touch. The immediacy of tactile experience has sometimes led touch to be thought of as privileged among senses and even infallible, as in Bishop Berkeley's famous claim that "touch tutors vision". By documenting novel ways in which touch goes wrong, this paper shows that we may need to think twice when we feel an insect land on our arm, or we may swat the wrong limb entirely. Good news if you're a mosquito.

## References

1. de Vignemont, F. (2018). *Mind the body: An exploration of bodily self-awareness* (Oxford: University Press).
2. Halligan, P.W., Hunt, M., Marshall, J.C., and Wade, D.T. (1995). Sensory detection without localization. *Neurocase 1*, 259–266.
3. Head, H., and Holmes, G. (1911). Sensory disturbances from cerebral lesions. *Brain 34*, 102–254.
4. Penfield, W., and Boldrey, E. (1937). Somatic motor and sensory representation in the cerebral cortex of man as studied by electrical stimulation. *Brain 60*, 389–443.
5. Nicolelis, M.A., Ghazanfar, A.A., Stambaugh, C.R., Oliveira, L.M., Laubach, M., Chapin, J.K., Nelson, R.J., and Kaas, J.H. (1998). Simultaneous encoding of tactile information by three primate cortical areas. *Nat. Neurosci. 1*, 621–630.
6. Grüsser, S.M., Mühlnickel, W., Schaefer, M., Villringer, K., Christmann, C., Koeppel, C., and Flor, H. (2004). Remote activation of referred phantom sensation and cortical reorganization in human upper extremity amputees. *Exp. Brain Res. 154*, 97–102.
7. Badde, S., Röder, B., and Heed, T. (2019). Feeling a touch to the hand on the foot. *Curr. Biol.*
8. Longo, M.R., Azañón, E., and Haggard, P. (2010). More than skin deep: body representation beyond primary somatosensory cortex. *Neuropsychologia 48*, 655–668.

9. Elbert, T., Pantev, C., Wienbruch, C., Rockstroh, B., and Taub, E. (1995). Increased cortical representation of the fingers of the left hand in string players. *Science* 270, 305–307.
10. Iwamura, Y., Iriki, A., and Tanaka, M. (1994). Bilateral hand representation in the postcentral somatosensory cortex. *Nature* 369, 554–556.
11. Tamè, L., Braun, C., Lingnau, A., Schwarzbach, J., Demarchi, G., Li Hegner, Y., Farnè, A., and Pavani, F. (2012). The contribution of primary and secondary somatosensory cortices to the representation of body parts and body sides: an fMRI adaptation study. *J. Cogn. Neurosci.* 24, 2306–2320.
12. Tamè, L., Azañón, E., and Longo, M.R. (2019). A conceptual model of tactile processing across body features of size, shape, side, and spatial location. *Front. Psychol.* 10, 291.
13. Pons, T.P., Garraghty, P.E., and Mishkin, M. (1998). Lesion-induced plasticity in the second somatosensory cortex of adult macaques. *Proc. Natl. Acad. Sci.* 85, 5279–5281.
14. Haggard, P., Kitadono, K., Press, C., and Taylor-Clarke, M. (2006). The brain's fingers and hands. *Exp. brain Res.* 172, 94–102.
15. Manser-Smith, K., Tamè, L., and Longo, M.R. (2018). Tactile confusions of the fingers and toes. *J. Exp. Psychol. Hum. Percept. Perform.* 44, 1727–1738.
16. Zhang, C.Y., Aflalo, T., Revechkis, B., Rosario, E.R., Ouellette, D., Pouratian, N., and Andersen, R.A. (2017). Partially mixed selectivity in human posterior parietal association cortex. *Neuron* 95, 697–708.e4.
17. Treisman, A. (1996). The binding problem. *Curr. Opin. Neurobiol.* 6, 171–178.
18. Azañón, E., Stenner, M.-P., Cardini, F., and Haggard, P. (2015). Dynamic tuning of tactile localization to body posture. *Curr. Biol.* 25, 512–517.



19. Kosslyn, S.M. (2006). You can play 20 questions with nature and win: categorical versus coordinate spatial relations as a case study. *Neuropsychologia* 44, 1519–1523.
20. Yamamoto, S., and Kitazawa, S. (2001). Reversal of subjective temporal order due to arm crossing. *Nat. Neurosci.* 4, 759–765.

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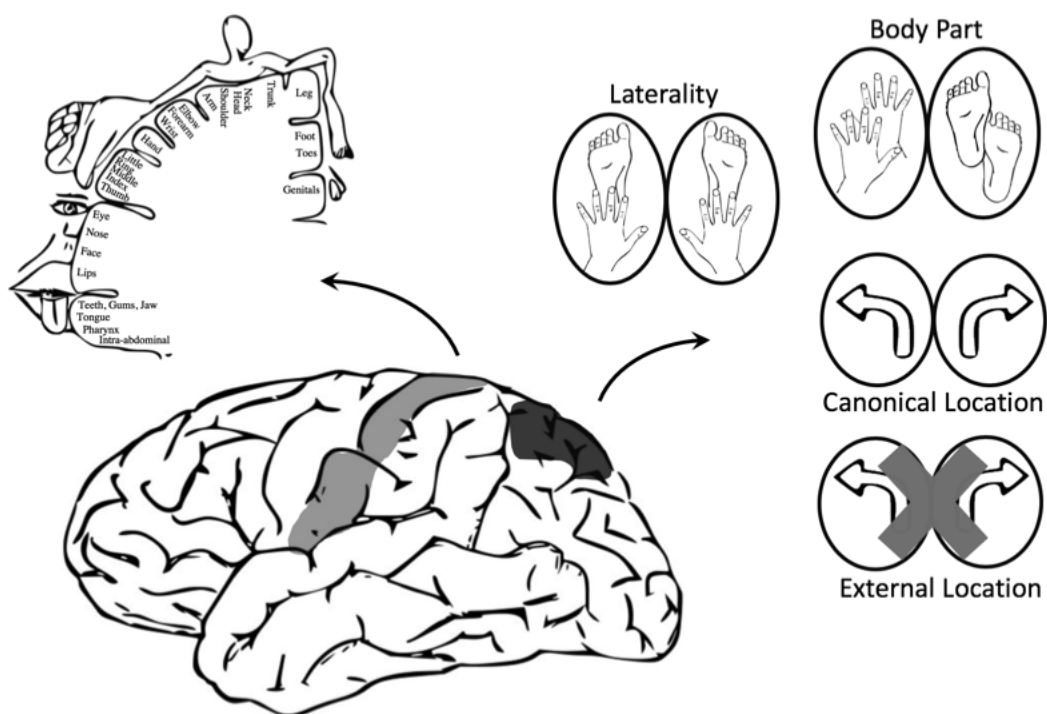
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**Figure 1. Continuous vs categorical representations in tactile localization.**

The ability to locate where on the continuous body surface touch has occurred putatively depends on the topographic maps observed in the primary somatosensory cortex (marked in soft grey). This is illustrated in what is known as the sensory homunculus, as an orderly topography of body parts in the cortex. Alternatively, the ability to locate touch on the skin might be based on abstract features or categorical information related to the touched body part. This second view is supported by Badde et al.'s [7] pattern of mislocalisations between

hands and feet, which are clearly not determined by proximity in the homunculus. These mislocalizations follow a systematic pattern consistent with the use of three main categorical features: 1) body part, by which a touch on the right hand might be misperceived to the left hand (i.e., the homologous limb); 2) body laterality, by which a touch on the right hand might be misperceived to the right foot; and 3) canonical side of the touched body, by which a touch on the crossed right hand might be misperceived to the foot placed at the right side, even though the touch was located at the left side of external space. Intriguingly, the external location of the touched limb did not predict the pattern of mislocalizations across hands and feet. The four types of categorical features are schematically represented on the right panel. We refer to posterior parietal brain regions (marked in dark grey) as the source for such putative representations, following evidence that these regions might represent categorical information, such as body parts and body sides in a partially mixed fashion [16].