

RUNNING HEAD: Vision and Implicit Hand Maps

The Effects of Immediate Vision on Implicit Hand Maps

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Word count: 3,472

Abstract

Perceiving the external spatial location of the limbs using position sense requires that

immediate proprioceptive afferent signals be combined with a stored body model specifying

the size and shape of the body. Longo and Haggard (2010) developed a method to isolate and

measure this body model in the case of the hand in which participants judge the perceived

location in external space of several landmarks on their occluded hand. The spatial layout of

judgments of different landmarks is used to construct implicit hand maps, which can then be

compared to actual hand shape. Studies using this paradigm have revealed that the body

model of the hand is massively distorted, in a highly stereotyped way across individuals, with

large underestimation of finger length and overestimation of hand width. Previous studies

using this paradigm have allowed participants to see the locations of their judgments on the

occluding board. Several previous studies have demonstrated that immediate vision, even

when wholly non-informative, can alter processing of somatosensory signals and alter the

reference frame in which they are localised. The present study therefore investigated whether

immediate vision contributes to the distortions of implicit hand maps described previously.

Participants judged the external spatial location of the tips and knuckles of their occluded left

hand either while being able to see where they were pointing (as in previous studies) or while

blindfolded. The characteristic distortions of implicit hand maps reported previously were

clearly apparent in both conditions, demonstrating that the distortions are not an artefact of

immediate vision. However, there were significant differences in the magnitude of distortions

in the two conditions, suggesting that vision may modulate representations of body size and

shape, even when entirely non-informative.

Keywords: Body Representation; Body Schema; Position Sense; Vision

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Several classes of afferent signals from the periphery provide information about the location of the limbs, including receptors in joints signalling flexion or extension, from the skin signalling stretch, and from muscle spindles signalling contraction or lengthening (Proske and Gandevia 2012). Each of these signals provides information body *posture*, that is, about the relative flexion or extension of each joint. In order to use these proprioceptive signals for position sense to localise the absolute spatial location of part of the body in external space, such information about joint angles must be combined with information about the size and shape of the body segments between joint. Critically, however, no afferent signals directly specify body size or shape, indicating that this information must come from a stored *body model* (Longo et al. 2010).

In a series of recent studies, my colleagues and I have investigated this body model using a novel 'psychomorphometric' method in which participants judge the perceived location of several landmarks on their occluded hand, and the pattern of responses is used to construct implicit perceptual maps of hand size and shape (Longo and Haggard 2010, 2012a, 2012b, Longo et al. 2012). In these studies, we have found that the body model of the hand is massively distorted, in a highly stereotyped way across people. Specifically, there were three clear patterns of distortions: (1) overestimation of hand width, quantified as the distance between pairs of knuckles; (2) overall underestimation of finger length, quantified as the distance between the knuckle and tip of each finger; and (3) a radio-ulnar gradient, with underestimation of finger length increasing systematically from the thumb to the little finger. In contrast, explicit estimates of hand shape were approximately veridical (Longo and Haggard 2010), suggesting that the body model is implicit, and outside of conscious bodily awareness.

The current paper focuses on one aspect of the procedure used in previous studies investigating implicit hand maps (e.g., Longo and Haggard 2010, 2012a, 2012b, Longo et al.

2012), namely the fact that participants have been able to see where they were pointing on each trial. Vision in this context is uninformative about the location of individual landmarks (except for a very broad sense in which seeing the boundaries of the occluding board imposes extreme limits on landmark location). Vision was allowed in these studies to ensure that participants' responses were as precise as possible, but was not considered to be an important contributing factor to the distortions observed. It is important to consider, however, whether immediate vision might be driving aspects of the results. Two recent studies investigating the effects of vestibular stimulation on body representation did have participants respond while blindfolded (Ferrè et al. 2013; Lopez et al. 2012). Ferrè and colleagues reported distortions similar to those reported by Longo and Haggard (2010). Lopez and colleagues did not report statistical comparisons of represented and actual hand size (focusing instead on vestibular stimulation versus sham), but the pattern of their results appears qualitatively consistent with distortions reported in other studies. In both these studies, participants were blindfolded in all conditions, making the potential effects of immediate vision impossible to estimate.

Several previous results suggest that vision, even when non-informative, affects several aspects of somatosensory processing. For example, in the *visual enhancement of touch* (Kennett et al. 2001; Cardini et al. 2011) and *visual analgesia* (Longo et al. 2009; Mancini et al. 2011) effects, vision of the stimulated body part enhances the spatial acuity of touch and reduces the perceived intensity of acute pain, respectively, even when completely non-informative about the stimulation. Those studies are importantly different from the proprioceptive localisation situation in that they involve vision of the body itself, rather than just in the direction of the body's location, but nevertheless show that non-informative vision can clearly affect somatosensation. Perhaps more directly relevant, Forster and Eimer (2005) found that looking in the direction of an unseen body part altered tactile attention on that

body part. These results clearly show that even when vision provides no actual task-relevant information, it may nevertheless be modulating somatosensory processing.

Further, numerous studies have suggested that vision can determine the reference frame for somatosensory perception. For example, Harris and colleagues have argued in a number of recent papers that tactile information is localised using a gaze-centred frame of reference (Harrar and Harris 2009, 2010; Pritchett and Harris 2011). Other results have suggested that when vision is present, kinaesthetic information may be automatically transformed into a visual reference frame (Darling and Miller 1993). Indeed, recent studies have shown effects of gaze direction on proprioceptive localisation judgments (Fiehler et al. 2010; Jones and Henriques 2010). Collectively, these results suggest profound interactions between vision and both somatosensation generally, and proprioception specifically. Thus, it cannot be assumed that the presence of immediate vision in previous studies of implicit hand maps has not contributed to the distortions observed.

The present study therefore compared implicit hand maps obtained using the method of Longo and Haggard (2010) in conditions in which participants could see where they were pointing on the occluding board (as in previous studies) and while blindfolded. To the extent that distortions observed in previous studies reflect transformation of proprioceptive information into a gaze-centred reference frame, they should not appear when participants respond while blindfolded. In contrast, to the extent that the distortions reflect characteristics of stable body representations underlying position sense, they should not be dependent on the presence of immediate vision.

Method

Participants

Twelve individuals (eight female) between 19 and 34 years of age from the University of London community participated for payment. All but one were right handed as assessed by the Edinburgh Inventory (M: 74.88; range: -100 to +100). Procedures were approved by the local ethics committee.

Procedures

Procedures were similar to those I have used in previous studies with this paradigm (e.g., Longo & Haggard, 2010, 2012a, 2012b; Longo et al., 2012). Participants sat with their left hand lying palm-down on a table, approximately aligned with their body midline. Participants were asked to sit upright, with their torso up against the edge of the table. The hand rested flat on the table, with fingers straight. An occluding board (40 x 40 cm) rested on four pillars (6 cm in height) and covered the hand. A webcam (Creative Live Cam Voice) suspended approximately 27 cm above the table captured still photographs (1280 x 960 pixels) under control of a custom MATLAB (Mathworks, Natick, MA) script.

In the *Sighted* condition, participants responded as in my previous studies using this paradigm, with full vision of the occluding board and the stick they were pointing with. In the *Blindfolded* condition, in contrast, participants wore a blindfold so that the task had to be completed using only haptic feedback.

Participants used a short wooden stick (11.4 cm in length) to indicate the perceived location of landmarks on their left hand. In my previous studies using this paradigm, participants used a long (35 cm) baton to point to different landmarks on their hand. Because participants in this study had to point without visual guidance, a shorter stick was used, so that participants could more easily feel where the tip was pointing. The tip of the stick tapered gradually to a point. Participants made judgments about ten different landmarks: the tip (i.e., the most distal point) and the knuckle (i.e., centre of the knuckle at the base of each

finger) of each finger. Participants were given verbal instructions on each trial about which landmark to judge. At the beginning of the experiment, they were instructed that they should be precise in their responses, avoid ballistic pointing, and avoid strategies such as tracing the outline of the hand. To ensure that responses on each trial were independent, participants were asked to move the stick to tap a metal post that rested to the side of the board, within easy reach of their right hand. The post was used in place of the blue dot used in previous studies since it could be found even when participants were blindfolded. A photograph of each response was taken when the participant indicated that they were happy with their response.

There were four blocks of thirty trials each. Sighted and blindfolded blocks were counterbalanced in an ABBA fashion, with the first condition counterbalanced across participants. Each block consisted of three mini-blocks of ten trials each, including one of each of the ten landmarks in random order. At the beginning and end of each block an image was taken without the occluding board, which allowed measurement of the true proportions of the participant's hand, and confirmation that the hand had not moved during the block. To facilitate coding, a black dot was made on each of the knuckles with a black pen. A ruler (10 cm in length) was affixed to the table and appeared in the pictures without the occluding board, allowing conversion between distances in pixels and cm.

Analysis

The camera used a wide-angle ('fisheye') lens, which induced some spatial distortion. This fisheye distortion was corrected using the Panotools plug-in (http://www.panotools.com/) for Adobe Photoshop CS2. The x-y pixel coordinates of each landmark on the photographs of the hands and of the corresponding judged locations were coded using ImageJ software (Abramoff et al. 2004). The length of different parts of the hand

was then calculated. The length of each finger was calculated as the distance between the knuckle and tip of each finger. Hand width was similarly quantified as the distance between pairs of knuckles, with the distance between the knuckles of the index and little fingers being used as an overall measure of hand width. For each distance, percent overestimation was calculated as $100 \times (Judged Length - Actual Length) / Actual Length$.

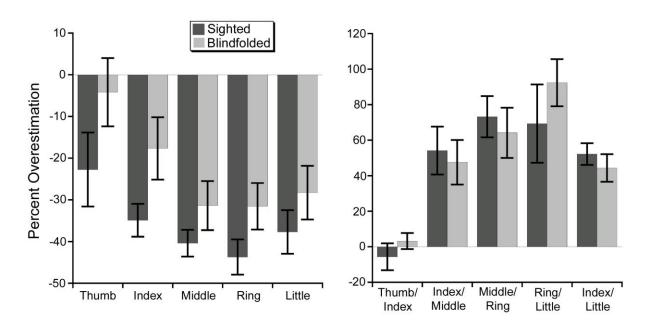
Maps of both actual and judged locations were put into Procrustes alignment using the CoordGen software, part of the Integrated Morphometric Program (IMP; David Sheets, Canisius College, http://www.canisius.edu/~sheets/morphsoft.html). Procrustes superimposition removes differences in location, rotation, and overall size to isolate differences in shape between configurations of homologous landmarks (Rohlf & Slice, 1990; Bookstein, 1991).

Because the fingers are articulated and can move independently, we rotated the fingertips of each map to be in a constant posture, as in our previous study (Longo & Haggard, 2010). First, we calculated the average angle of each finger relative to the hand in the photos of each participant's actual hands, defined as the angle between a line running through the knuckles of the index and little fingers and another line running through the knuckle and tip of each finger. These angles were 39.6°, 64.4°, 76.5°, 87.1°, and 108.8°, for the thumb through little fingers, respectively. Then for each experimental block and for each participant, the fingertip of each finger was rotated so that the fingers were at those angles (i.e., hand posture was matched across blocks and across participants).

As there were two experimental blocks for each task, maps from each participant were first put into Procrustes alignment so that an average map could be calculated for each participant for both actual location and judgments in each of the two tasks. Then a second-level generalized Procrustes analysis was conducted putting actual and judged maps from all participants into simultaneous alignment, separately for the two tasks.

Results

All three of the patterns of distortion I have reported in my previous studies using this paradigm (Longo & Haggard, 2010, 2012a, 2012b) were investigated here. The left panel of Figure 1 shows overestimation of finger length. The values are negative, indicating overall underestimation. Averaging across the five fingers, there was clear underestimation of finger length in both the sighted (-35.9% underestimation), t(11) = -7.73, p < .0001, and blindfolded (-22.6% underestimation), t(11) = -3.85, p < .005, conditions. Nevertheless, there was a significant reduction in the magnitude of underestimation in the blindfolded condition, t(11) = 2.24, p < .05. There was a modest (but non-significant) positive correlation between the magnitude of underestimation in the two conditions, t(10) = 0.381, t = 0.3

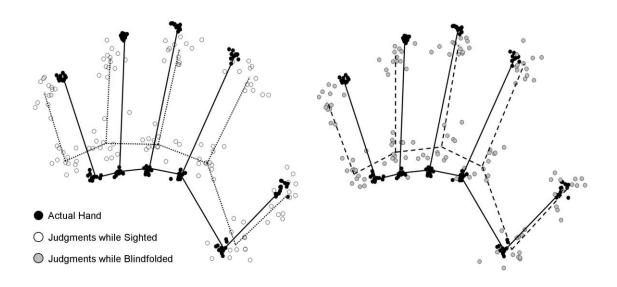


<u>Figure 1</u>: Left panel: Percent overestimation (i.e., 100*(Judged Length – Actual Length)/Actual Length) for each of the five fingers in the two conditions. *Right panel*: Percent overestimation for the distance between pairs of knuckles. The distance between the index and little fingers is taken as an overall measure of hand width. Error bars are one S.E.M.

To assess the gradient of underestimation across fingers, least-squares regression was used to estimate the increase in underestimation with a shift of one digit towards the little

finger. Consistent with previous findings, there was a clear increase in underestimation from the thumb to little finger in both the sighted (mean β = 3.9% / finger), t(11) = 2.97, p < .02, and blindfolded (mean β = 6.2% / finger), t(11) = 4.92, p < .001, conditions. The magnitude of this gradient was marginally larger in the blindfolded condition, t(11) = 2.01, p = .069. There was a significant correlation between slopes in the two conditions, r(10) = 0.594, p < .05.

The right panel of Figure 1 shows overestimation of the distance between pairs of knuckles. The distance between the knuckles of the index and little fingers was taken as an overall measure of hand width. There was clear overestimation of hand width in both the sighted (52.2% overestimation), t(11) = 8.54, p < .0001, and blindfolded (44.3% overestimation), t(11) = 5.70, p < .0001, conditions. The magnitude of overestimation did not differ significantly between the two conditions, t(11) = 1.11, n.s. There was a marginally significant correlation between the magnitude of overestimation in the two conditions, t(10) = 0.508, t=0.092 (two-tailed).



<u>Figure 2</u>: Hand maps in the two conditions placed in generalised Procrustes superimposition (GPS) alignment with maps of each participant's actual hand.

The overall aspect ratio of the hand was quantified using Napier's (1980) *shape index*, a ratio of hand width to length. Following my previous studies (Longo & Haggard, 2010, 2012b), the distance between the knuckles of the index and little fingers was taken as a measure of hand width, and the length of the middle finger as a measure of hand length. The shape index is calculated as: SI = 100 x (width / length). Large shape indices indicate a squat, wide hand, while small values indicate a thin, slender hand. The shape index was calculated for each participant for their actual hand, as well as for the hand maps in each condition (shown in the left 3 bars of Figure 3). The shape indices for hand maps were significantly greater than for the actual hand in both the sighted (150.8 vs. 58.0), t(11) = 12.04, p < .0001, and blindfolded (129.6 vs. 58.0), t(11) = 6.87, p < .0001, conditions. The shape indices for the two experimental conditions did not differ significantly from each other, t(11) = 1.50, n.s.

One interesting difference between the present study and previous studies using the same paradigm (e.g., Longo & Haggard, 2010, 2012a, 2012b) is the use of a much shorter stick for responding in this study (11.4 vs. 35.0 cm). In order to investigate any potential effect of this variable, the right six bars of Figure 3 show the shape indices from six previous experiments which were similar to the sighted condition of this study except for the length of the stick. These experiments were: (1) Exp. 1 from Longo and Haggard (2010); (2) the 'normal' posture from Exp. 2 from Longo and Haggard (2010); (3) the left hand condition from Exp. 3 from Longo and Haggard (2010); (4) Exp. 4 from Longo and Haggard (2010); (5) the dorsum condition from Longo and Haggard (2012a); (6) the 'localisation' task from Longo and Haggard (2012b). As is clear from Figure 3, the shape indices in the present study are similar to those found in previous studies using a longer baton, suggesting that the length of the implement used for responses does not have a major effect on the results.

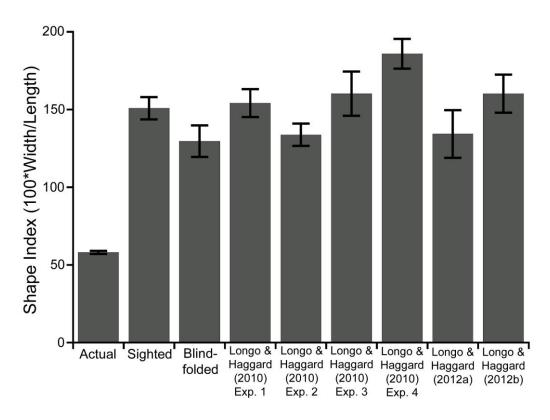


Figure 3: The three leftmost bars show shape indices (i.e., 100xWidth/Length) for participants' actual hands and for hand maps in the sighted and blindfolded conditions. Shape indices were clearly larger for the hand maps than for actual hand shape. For comparison, the rightmost six bars present the same values from six previous experiments using the same paradigm with participants responding using a long baton (35 cm). Error bars are one S.E.M.

Discussion

Similar patterns of distortion of implicit body representations underlying position sense were found regardless of whether participants had immediate vision while making their responses or not. Three types of distortions of implicit hand maps were apparent in both conditions, replicating recent results (Longo and Haggard 2010, 2012a, 2012b, Longo et al. 2012): (1) overall underestimation of finger length, (2) increasing underestimation of length from thumb to little finger, and (3) overall overestimation of hand width. These results demonstrate that the patterns of distortion we have previously reported are not dependent on immediate vision, and are unlikely to reflect an artefact of responses being coded in a visual reference frame.

These results have implications for understanding the origins of the distortions we have observed. One critical question concerns whether they reflect distortions representations of the body, per se, or whether they emerge from the manner in which participants are asked to respond. The present results contribute to a growing body of evidence that the distortions we have reported show a high level of generality across different modes of response, reducing the likelihood that they reflect an artefact of any specific mode of response. Similar distorted maps of the hand are apparent when participants have both visual and proprioceptive feedback about the location of their responses (Longo & Haggard, 2010; this study), when they have only proprioceptive feedback about their responding hand (this study), and when they have only visual feedback about their responses (i.e., they verbally guided a naïve experimenter who held the baton, Longo et al. 2012). Further, we found different patterns of distortions for judgments about two sides of a single body part (i.e., the dorsal and palmar hand surfaces, Longo and Haggard 2012a), though the manner in which responses were made was identical in both cases. Together, these results suggest that the measured distortions reflect distorted representations of the body itself, rather than idiosyncratic aspects of the manner in which participants indicate their judgments.

While qualitatively similar patterns of distortion were observed in both the sighted and blindfolded conditions, there was evidence that the magnitude of these distortions differed between them. These results are consistent with the possibility that vision, even though non-informative about the location of the body, might nevertheless alter somatosensory processing (cf. Kennett et al. 2001; Forster & Eimer 2005) or change the reference frame for somatosensory spatial perception (cf. Fiehler et al. 2010; Harrar and Harris 2009). Immediate vision led to a significant reduction in perceived finger length. The direction of this effect is consistent with the recent finding that non-informative vision of the hand leads to a reduction in the perceived tactile distance on the hand (Longo and Sadibolova

2013). The reason for vision producing reductions in represented body part size in both these cases are not fully clear. One possibility is that visually-induced reduction may be the converse of the perceived swelling seen following acute deafferentation (Gandevia and Phegan 1999) and in chronic pain (Moseley 2005; Pelz et al. 2011). Both deafferentation (Calford and Tweedale 1991) and chronic pain (Schwenkreis et al. 2003) are known to be associated with disinhibition of sensorimotor cortex. As suggested by Haggard, Iannetti, and Longo (2013), by increasing such inhibition (Cardini et al. 2011), vision may produce a constellation of effects opposite to those seen in chronic pain, including analgesia (Longo et al. 2009; Mancini et al. 2011), heightened tactile spatial acuity (Kennett et al. 2001; Cardini et al. 2011), and reduced represented body part size (Longo and Sadibolova 2013; this study).

Such an interpretation treats the blindfolded condition as a baseline, reflecting simply the absence of vision. Another possibility, however, is that blindfolding may induce rapid plasticity of somatosensory representations, as has been shown by several previous results (Facchini and Aglioti 2003; Weisser et al. 2005; Wong et al. 2001; Zubek et al. 1964). Indeed, Wong and colleagues (2001) reported changes in tactile spatial acuity after periods of blindfolding as short as 10 minutes, similar to the length of time participants in the present study were blindfolded during a single block of trials. Thus, it is also possible to consider the sighted condition as the baseline, and the differences observed in the blindfolded condition as reflecting short-term plasticity of body representations induced by blindfolding. It is difficult to see what type of condition would be a true baseline in this context to determine whether the differences seen between conditions reflect changes induced by immediate vision or short-term plasticity induced by blindfolding. This is an interesting question for future research.

References

- Abramoff MD, Magelhaes PJ, Ram SJ (2004) Image processing with ImageJ. Biophotonics Intern 11: 36-42
- Bookstein FL (1991) Morphometric tools for landmark data. Cambridge: Cambridge University Press.
- Calford MB, Tweedale R (1991) C-fibres provide a source of masking inhibition to primary somatosensory cortex. Proc Roy Soc B 243: 269-275
- Cardini F, Longo MR, Haggard P (2011) Vision of the body modulates somatosensory intracortial inhibition. Cereb Cortex 21: 2014-2022
- Cardini F, Longo MR, Driver J, Haggard P (2012) Rapid enhancement of touch from non-informative vision of the hand. Neuropsychologia 50: 1954-1960
- Darling WG, Miller GF (1993) Transformations between visual and kinaesthetic coordinate systems in reaches to remembered object locations and orientations. Exp Brain Res 93: 534-547
- Facchini S, Aglioti SM (2003) Short term light deprivation increases tactile spatial acuity in humans. Neurol 60: 1998-1999
- Ferrè ER, Vagnoni E, Haggard P (2013) Vestibular contributions to bodily awareness.

 Neuropsychologia 51: 1445-1452
- Fiehler K, Rösler F, Henriques DY (2010) Interaction between gaze and visual and proprioceptive position judgments. Exp Brain Res 203: 485-498
- Forster B, Eimer M (2005) Vision and gaze direction modulate tactile processing in somatosensory cortex: evidence from event-related brain potentials. Exp Brain Res 165: 8-18
- Gandevia SC, Phegan CM (1999) Perceptual distortions of the human body image produces by local anesthesia, pain and cutaneous stimulation. J Physiol 514: 609-616

- Haggard P, Iannetti GD, Longo MR (2013) Spatial sensory organization and body representation in pain perception. Curr Biol 23: R164-R176
- Harrar V, Harris LR (2009) Eye position affects the perceived position of touch. Exp Brain Res 198: 403-410
- Harrar V, Harris LR (2010) Touch used to guide action is partially coded in a visual reference frame. Exp Brain Res 203: 615-620
- Harrar V, Pritchett LM, Harris LR (2013) Segmented space: Measuring tactile localisation in body coordinates. Multisens Res 26: 3-18
- Jones SA, Henriques DY (2010) Memory for proprioceptive and multisensory targets is partially coded relative to gaze. Neuropsychologia 48: 3782-3792
- Kennett S, Taylor-Clarke M, Haggard P (2001) Noninformative vision improves the spatial resolution of touch in humans. Curr Biol 11: 1188-1191
- Leon-Sarmiento FE, Hernandez HG, Schroeder N (2008) Abnormal tactile discrimination and somatosensory plasticity in familial primary hyperhidrosis. Neurosci Lett 441: 332-334
- Longo MR, Haggard P (2010) An implicit body representation underlying human position sense. Proc Ntnl Acad Sci USA 107: 11727-11732
- Longo MR, Haggard P (2012a) A 2.5-D representation of the human hand. J Exp Psychol: Human Percept Perform 38: 9-13
- Longo MR, Haggard P (2012b) Implicit body representations and the conscious body image. Acta Psychol 141: 164-168
- Longo MR, Sadibolova R (2013) Seeing the body distorts tactile size perception. Cognition 126: 475-481
- Longo MR, Azañón E, Haggard P (2010). More than skin deep: Body representation beyond primary somatosensory cortex. Neuropsychologia 48: 655-668

- Longo MR, Betti V, Aglioti SM, Haggard P (2009) Visually induced analgesia: Seeing the body reduces acute pain. J Neurosci 29: 12125-12130
- Longo MR, Long C, Haggard P (2012) Mapping the invisible hand: A body model of a phantom limb. Psychol Sci 23: 740-742
- Lopez C, Schreyer H-M, Preuss N, Mast FW (2012) Vestibular stimulation modifies the body schema. Neuropsychologia 50: 1830-1837
- Mancini F, Longo MR, Kammers MPM, Haggard P (2011) Visual distortion of body size modulate pain perception. Psychol Sci 22: 325-330
- Moseley GL (2005) Distorted body image in complex regional pain syndrome. Neurology 65: 773
- Napier J (1980) Hands. Princeton: Princeton University Press.
- Pelz E, Seifert F, Lanz S, Müller R, Maihöfner C (2011) Impaired hand size estimation in CRPS. J Pain 12: 1095-1101
- Pritchett LM, Harris LR (2011) Perceived touch location is coded using a gaze signal. Exp Brain Res 213: 229-234
- Proske U, Gandevia SC (2012) The proprioceptive senses: Their roles in signalling body shape, body position and movement, and muscle force. Physiol Revs 92: 1651-1697
- Rholf FJ, Slice DE (1990) Extensions of the Procrustes method for the optimal superimposition of landmarks. Syst Zool 39: 40-59
- Schwenkreis P, Janssen F, Rommel O, Pleger B, Völker B, Hosbach I, Dertwinkel R, Maier C, Tegenthoff M (2003) Bilateral motor cortex disinhibition in complex regional pain syndrome (CRPS) type I of the hand. Neurology 61: 515-519
- Weisser V, Stilla R, Peltier S, Hu X, Sathian K (2005) Short-term visual deprivation alters neural processing of tactile form. Exp Brain Res 166: 572-582
- Wong M, Hackeman E, Hurd C, Goldreich D (2011) Short-term visual deprivation does not

enhance passive tactile spatial acuity. PLOS ONE 6: e25277

Zubek JP, Flye J, Aftanas M (1964) Cutaneous sensitivity after prolonged visual deprivation.

Science 144: 1591-1593

Acknowledgments

This research was supported by a grant from the European Research Council (ERC-2013-StG-336050) to MRL.